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(54) Title: METHODS AND COMPOSITIONS FOR TREATING HEPATITIS C VIRUS

METHODS AND COMPOSITIONS FOR TREATING HEPATITIS C VIRUS

FIELD OF THE INVENTION

This invention is in the area of pharmaceutical chemistry, and is in particular, is a compound, method and composition for the treatment of hepatitis C virus. This application claims priority to U.S. provisional application no. 60/206,585, filed on May 23, 2000.

BACKGROUND OF THE INVENTION

The hepatitis C virus (HCV) is the leading cause of chronic liver disease worldwide. (Boyer, N. et al. J. Hepatol. 32:98-112, 2000). HCV causes a slow growing viral infection and is the major cause of cirrhosis and hepatocellular carcinoma (Di Besceglie, A. M. and Bacon, B. R., Scientific American, Oct.: 80-85, (1999); Boyer, N. et al. J. Hepatol. 32:98-112, 2000). An estimated 170 million persons are infected with HCV worldwide. (Boyer, N. et al. J. Hepatol. 32:98-112, 2000). Cirrhosis caused by chronic hepatitis C infection accounts for 8,000-12,000 deaths per year in the United States, and HCV infection is the leading indication for liver transplant.

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HCV is known to cause at least 80% of posttransfusion hepatitis and a substantial proportion of sporadic acute hepatitis. Preliminary evidence also implicates HCV in many cases of "idiopathic" chronic hepatitis, "cryptogenic" cirrhosis, and probably hepatocellular carcinoma unrelated to other hepatitis viruses, such as Hepatitis B Virus (HBV). A small proportion of healthy persons appear to be chronic HCV carriers, varying with geography and other epidemiological factors. The numbers may substantially exceed those for HBV, though information is still preliminary; how many of these persons have subclinical chronic liver disease is unclear. (The Merck Manual, ch. 69, p. 901, 16th ed., (1992)).

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HCV has been classified as a member of the virus family Flaviviridae that includes the *genera* flaviviruses, pestiviruses, and hapaceiviruses which includes hepatitis C viruses (Rice, C. M., Flaviviridae: The viruses and their replication. *In*: Fields Virology, Editors: Fields, B. N., Knipe, D. M., and Howley, P. M., Lippincott-Raven Publishers, Philadelphia, PA, Chapter 30, 931-959, 1996). HCV is an enveloped virus containing a positive-sense single-stranded RNA genome of approximately 9.4kb. The viral genome consists of a 5' untranslated region (UTR), a long open reading frame encoding a polyprotein precursor of

approximately 3011 amino acids, and a short 3' UTR. The 5' UTR is the most highly conserved part of the HCV genome and is important for the initiation and control of polyprotein translation. Translation of the HCV genome is initiated by a cap-independent mechanism known as internal ribosome entry. This mechanism involves the binding of ribosomes to an RNA sequence known as the internal ribosome entry site (IRES). An RNA pseudoknot structure has recently been determined to be an essential structural element of the HCV IRES. Viral structural proteins include a nucleocapsid core protein (C) and two envelope glycoproteins, E1 and E2. HCV also encodes two proteinases, a zinc-dependent metalloproteinase encoded by the NS2-NS3 region and a serine proteinase encoded in the NS3 region. These proteinases are required for cleavage of specific regions of the precursor polyprotein into mature peptides. The carboxyl half of nonstructural protein 5, NS5B, contains the RNA-dependent RNA polymerase. The function of the remaining nonstructural proteins, NS4A and NS4B, and that of NS5A (the amino-terminal half of nonstructural protein 5) remain unknown.

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A significant focus of current antiviral research is directed toward the development of improved methods of treatment of chronic HCV infections in humans (Di Besceglie, A. M. and Bacon, B. R., *Scientific American*, Oct.: 80-85, (1999)). Currently, there are two primary antiviral compounds, Ribavirin and interferon-alpha, which are used for the treatment of chronic HCV infections in humans.

Treatment of HCV Infection with Ribivarin

Ribavirin (1- β -D-ribofuranosyl-1-1,2,4-triazole-3-carboxamide) is a synthetic, non-interferon-inducing, broad spectrum antiviral nucleoside analog sold under the trade name, Virazole (The Merck Index, 11th edition, Editor: Budavari, S., Merck & Co., Inc., Rahway, NJ, p1304, 1989). United States Patent No. 3,798,209 and RE29,835 disclose and claim Ribavirin. Ribavirin is structurally similar to guanosine, and has in vitro activity against several DNA and RNA viruses including *Flaviviridae* (Gary L. Davis. *Gastroenterology* 118:S104-S114, 2000).

Ribavirin reduces serum amino transferase levels to normal in 40% or patients, but it does not lower serum levels of HCV-RNA (Gary L. Davis. *Gastroenterology* 118:S104-S114, 2000). Thus, Ribavirin alone is not effective in reducing viral RNA levels. Additionally, Ribavirin has significant toxicity and is known to induce anemia.

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Treatment of HCV Infection with Interferon

Interferons (IFNs) are compounds that have been commercially available for the treatment of chronic hepatitis for nearly a decade. IFNs are glycoproteins produced by immune cells in response to viral infection. IFNs inhibit viral replication of many viruses, including HCV, and when used as the sole treatment for hepatitis C infection, IFN suppresses serum HCV-RNA to undetectable levels. Additionally, IFN normalizes serum amino transferase levels. Unfortunately, the effects of IFN are temporary and a sustained response occurs in only 8%-9% of patients chronically infected with HCV (Gary L. Davis. Gastroenterology 118:S104-S114, 2000).

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A number of patents disclose HCV treatments using interferon-based therapies. For example, U.S. Patent No. 5,980,884 to Blatt et al. discloses methods for retreatment of patients afflicted with HCV using consensus interferon. U.S. Patent No. 5,942,223 to Bazer et al. discloses an anti-HCV therapy using ovine or bovine interferon-tau. U.S. Patent No. 5,928,636 to Alber et al. discloses the combination therapy of interleukin-12 and interferon alpha for the treatment of infectious diseases including HCV. U.S. Patent No. 5,908,621 to Glue et al. discloses the use of polyethylene glycol modified interferon for the treatment of HCV. U.S. Patent No. 5,849,696 to Chretien et al. discloses the use of thymosins, alone or in combination with interferon, for treating HCV. U.S. Patent No. 5,830,455 to Valtuena et al. discloses a combination HCV therapy employing interferon and a free radical scavenger. U.S. Patent No. 5,738,845 to Imakawa discloses the use of human interferon tau proteins for treating HCV. Other interferon-based treatments for HCV are disclosed in U.S. Patent No. 5,676,942 to Testa et al., U.S. Patent No. 5,372,808 to Blatt et al., and U.S. Patent No. 5,849,696.

Combination of Interferon and Ribavirin

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The combination of IFN and Ribavirin for the treatment of HCV infection has been reported to be effective in the treatment of IFN naïve patients (Battaglia, A.M. et al., Ann. Pharmacother. 34:487-494, 2000). Results are promising for this combination treatment both before hepatitis develops or when histological disease is present (Berenguer, M. et al. Antivir. Ther. 3(Suppl. 3):125-136, 1998). Side effects of combination therapy include

hemolysis, flu-like symptoms, anemia, and fatigue. (Gary L. Davis. *Gastroenterology* 118:S104-S114, 2000).

Additional References Disclosing Methods to Treat HCV Infections

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A number of HCV treatments are reviewed by Bymock et al. in Antiviral Chemistry & Chemotherapy, 11:2; 79-95 (2000).

Several substrate-based NS3 protease inhibitors have been identified in the literature, in which the scissile amide bond of a cleaved substrate is replaced by an electrophile, which interacts with the catalytic serine. Attwood et al. (1998) Antiviral peptide derivatives, 98/22496; Attwood et al. (1999), Antiviral Chemistry and Chemotherapy 10.259-273; Attwood et al. (1999) Preparation and use of amino acid derivatives as anti-viral agents, German Patent Publication DE 19914474; Tung et al. (1998) Inhibitors of serine proteases, particularly hepatitis C virus NS3 protease, WO 98/17679. The reported inhibitors terminate in an electrophile such as a boronic acid or phosphonate. Llinas-Brunet et al. (1999) Hepatitis C inhibitor peptide analogues, WO 99/07734. Two classes of electrophile-based inhibitors have been described, alphaketoamides and hydrazinoureas.

The literature has also described a number of non-substrate-based inhibitors. For example, evaluation of the inhibitory effects of 2,4,6-trihydroxy-3-nitro-benzamide derivatives against HCV protease and other serine proteases has been reported. Sudo, K. et al., (1997) Biochemical and Biophysical Research Communications, 238:643-647; Sudo, K. et al. (1998) Antiviral Chemistry and Chemotherapy 9:186. Using a reverse-phase HPLC assay, the two most potent compounds identified were RD3-4082 and RD3-4078, the former substituted on the amide with a 14 carbon chain and the latter processing a paraphenoxyphenyl group.

Thiazolidine derivatives have been identified as micromolar inhibitors, using a

reverse-phase HPLC assay with an NS3/4A fusion protein and NS5A/5B substrate. Sudo, K. et al. (1996) Antiviral Research 32:9-18. Compound RD-1-6250, possessing a fused cinnamoyl moiety substituted with a long alkyl chain, was the most potent against the

isolated enzyme. Two other active examples were RD4 6205 and RD4 6193.

Other literature reports screening of a relatively small library using an ELISA assay and the identification of three compounds as potent inhibitors, a thiazolidine and two benzanilides. Kakiuchi N. et al. J. EBS Letters 421:217-220; Takeshita N. et al., Analytical Biochemistry 247:242-246, 1997. Several U.S. patents disclose protease inhibitors for the treatment of HCV. For example, U.S. Patent No. 6,004,933 to Spruce et al. discloses a class of cysteine protease inhibitors for inhibiting HCV endopeptidase 2. U.S. Patent No. 5,990,276 to Zhang et al. discloses synthetic inhibitors of hepatitis C virus NS3 protease. The inhibitor is a subsequence of a substrate of the NS3 protease or a substrate of the NS4A cofactor. The use of restriction enzymes to treat HCV is disclosed in U.S. Patent No. 5,538,865 to Reyes et al.

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Isolated from the fermentation culture broth of *Streptomyces* sp., Sch 68631, a phenan-threnequinone, possessed micromolar activity against HCV protease in a SDS-PAGE and autoradiography assay. Chu M. *et al.*, *Tetrahedron Letters* 37:7229-7232, 1996. In another example by the same authors, Sch 351633, isolated from the fungus *Penicillium griscofuluum*, demonstrated micromolar activity in a scintillation proximity assay. Chu M. *et al.*, *Bioorganic and Medicinal Chemistry Letters* 9:1949-1952. Nanomolar potency against the HCV NS3 protease enzyme has been achieved by the design of selective inhibitors based on the macromolecule eglin c. Eglin c, isolated from leech, is a potent inhibitor of several serine proteases such as S. griseus proteases A and B, α-chymotrypsin, chymase and subtilisin. Qasim M.A. *et al.*, *Biochemistry* 36:1598-1607, 1997.

HCV helicase inhibitors have also been reported. U.S. Patent No. 5,633,358 to Diana G.D. et al.; PCT Publication No. WO 97/36554 of Diana G.D. et al.. There are a few reports of HCV polymerase inhibitors: some nucleotide analogues, gliotoxin and the natural product cerulenin. Ferrari R. et al., Journal of Virology 73:1649-1654, 1999; Lohmann V. et al., Virology 249:108-118, 1998.

Antisense phosphorothioate oligodeoxynucleotides complementary to sequence stretches in the 5' non-coding region of the HCV, are reported as efficient inhibitors of HCV gene expression in *in vitro* translation and IIcpG2 IICV-luciferase cell culture systems. Alt M. *et al.*, *Hepatology* 22:707-717, 1995. Recent work has demonstrated that nucleotides 326-348 comprising the 3' end of the NCR and nucleotides 371-388 located in the core coding region of the HCV RNA are effective targets for antisense-mediated inhibition of viral translation. Alt M. *et al.*, *Archives of Virology* 142:589-599, 1997. U.S.

Patent No. 6,001,990 to Wands et al. discloses oligonucleotides for inhibiting the replication of HCV. PCT Publication No. WO 99/29350 discloses compositions and methods of treatment for hepatitis C infection comprising the administration of antisense oligonucleotides that are complementary and hybridizable to HCV-RNA. U.S. Patent No. 5,922,857 to Han et al. disclose nucleic acids corresponding to the sequence of the pestivirus homology box IV area for controlling the translation of HCV. Antisense oligonucleotides as therapeutic agents have been recently reviewed (Galderisi U. et al., Journal of Cellular Physiology 181:251-257, 1999).

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Other compounds have been reported as inhibitors of IRES-dependent translation in HCV. Japanese Patent Publication JP-08268890 of Ikeda N et al.; Japanese Patent Publication JP-10101591 of Kai, Y. et al. Nuclease-resistant ribozymes have been targeted at the IRES and recently reported as inhibitors in an HCV-poliovirus chimera plaque assay. Maccjak D.J. et al., Hepatology 30 abstract 995, 1999. The use of ribozymes to treat HCV is also disclosed in U.S. Patent No. 6,043,077 to Barber et al., and U.S. Patent Nos. 5,869,253 and 5,610,054 to Draper et al.

Other patents disclose the use of immune system potentiating compounds for the treatment of HCV. For example, U.S. Patent No. 6,001,799 to Chretien *et al.* discloses a method of treating hepatitis C in non-responders to interferon treatment by administering an immune system potentiating dose of thymosin or a thymosin fragment. U.S. Patent Nos. 5,972,347 to Eder *et al.* and 5,969,109 to Bona *et al.* disclose antibody-based treatments for treating HCV.

U.S. Patent No. 6,034,134 to Gold *et al.* discloses certain NMDA receptor agonists having immunodulatory, antimalarial, anti-Borna virus and anti-Hepatitis C activities. The disclosed NMDA receptor agonists belong to a family of 1-amino-alkylcyclohexanes. U.S. Patent No. 6,030,960 to Morris-Natschke *et al.* discloses the use of certain alkyl lipids to inhibit the production of hepatitis-induced antigens, including those produced by the HCV virus. U.S. Patent No. 5,922,757 to Chojkier *et al.* discloses the use of vitamin E and other antioxidants to treat hepatic disorders including HCV. U.S. Patent No. 5,858,389 to Elsherbi *et al.* discloses the use of squalene for treating hepatitis C. U.S. Patent No. 5,849,800 to Smith et al discloses the use of amantadine for treatment of Hepatitis C. U.S. Patent No. 5,846,964 to Ozeki *et al.* discloses the use of bile acids for treating HCV. U.S.

Patent No. 5,491,135 to Blough *et al.* discloses the use of N-(phosphonoacetyl)-L-aspartic acid to treat flaviviruses such as HCV.

Other compounds proposed for treating HCV include plant extracts (U.S. Patent No. 5,837,257 to Tsai *et al.*, U.S. Patent No. 5,725,859 to Omer *et al.*, and U.S. Patent No. 6,056,961), piperidenes (U.S. Patent No. 5,830,905 to Diana *et al.*), benzenedicarboxamides (U.S. Patent No. 5,633,388 to Diana *et al.*), polyadenylic acid derivatives (U.S. Patent No. 5,496,546 to Wang *et al.*), 2',3'-dideoxyinosine (U.S. Patent No. 5,026,687 to Yarchoan *et al.*), benzimidazoles (U.S. Patent No. 5,891,874 to Colacino *et al.*).

In light of the fact that the hepatitis C virus has reached epidemic levels worldwide, and has tragic effects on the infected patient, there remains a strong need to provide new effective pharmaceutical agents to treat hepatitis C that has low toxicity to the host.

Therefore, it is an object of the present invention to provide a compound, method and composition for the treatment of a host infected with hepatitis C virus.

SUMMARY OF THE INVENTION

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Compounds, methods and compositions for the treatment of hepatitis C infection are described that include an effective hepatitis C treatment amount of a β -D- or β -L-nucleoside of the Formulas (I) – (XVIII), or a pharmaceutically acceptable salt or prodrug thereof.

In a first principal embodiment, a compound of Formula I, or a pharmaceutically acceptable salt or produce thereof, is provided:

$$X^1$$
 N
 N
 N
 X^2
 CH_3
 CH_3
 (I)

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wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl);

sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a second principal embodiment, a compound of Formula II, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

$$X^1$$
 N
 X^2
 N
 N
 X^2
 N
 N
 X^2
 N
 N
 X^2
 N
 N
 X
 X

wherein:

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 R^1 , R^2 and R^3 are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 or R^3 is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR4, NR4R5 or SR4;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a third principal embodiment, a compound of Formula III, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

$$X^1$$
 X^1
 X^2
 X^2
 X^2
 X^2
 X^2
 X^2
 X^2
 X^2
 X^3
 X^2
 X^3
 X^4
 X^2
 X^3
 X^4
 X^2

10 wherein:

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 R^1 , R^2 and R^3 are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a `peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 or R^3 is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

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In a fourth principal embodiment, a compound of Formula IV, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

5 wherein:

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R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a fifth principal embodiment, a compound of Formula V, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

$$X^{1}$$
 X^{1}
 X^{1

wherein:

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R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a sixth principal embodiment, a compound of Formula VI, or a pharmaceutically acceptable salt or produce thereof, is provided:

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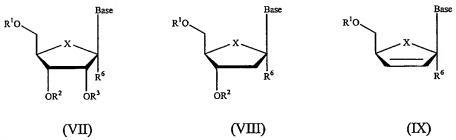
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R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a seventh principal embodiment, a compound selected from Formulas VII, VIII and IX, or a pharmaceutically acceptable salt or prodrug thereof, is provided:



wherein:

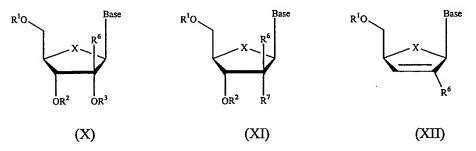
20 Base is a purine or pyrimidine base as defined herein;

 R^1 , R^2 and R^3 are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 or R^3 is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and

5 X is O, S, SO₂ or CH₂.

In a eighth principal embodiment, a compound of Formulas X, XI and XII, or a pharmaceutically acceptable salt or prodrug thereof, is provided:



wherein:

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Base is a purine or pyrimidine base as defined herein;

 R^1 , R^2 and R^3 are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 or R^3 is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(lower alkyl), -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(acyl)₂; and

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In a ninth principal embodiment a compound selected from Formulas XIII, XIV and XV, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

wherein:

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Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -NH(acyl), -N(acyl), -N(acyl), and

20 X is O, S, SO_2 or CH_2 .

In a tenth principal embodiment the invention provides a compound of Formula XVI, or a pharmaceutically acceptable salt or prodrug thereof:

R O Base
$$R^{10}$$
 R^{10} R^{8} R^{6} R^{9} R^{7} R^{6} (XVI)

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Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ or R² is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a pi bond; and

X is O, S, SO₂ or CH₂.

In a eleventh principal embodiment the invention provides a compound of Formula XVII, or a pharmaceutically acceptable salt or prodrug thereof:

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WO 01/90121 PCT/US01/16671 wherein:

Base is a purine or pyrimidine base as defined herein;

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R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ or R² is independently H or phosphate;

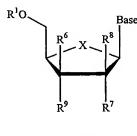
R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a pi bond; and X is O, S, SO₂ or CH₂.

In an twelfth principal embodiment, the invention provides a compound of Formula XVIII, or a pharmaceutically acceptable salt or prodrug thereof:



25 (XVIII)

WO 01/90121 wherein:

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PCT/US01/16671

Base is a purine or pyrimidine base as defined herein;

R¹ and R² independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ or R² is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(lower-alkyl)amino;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a pi bond;

20 X is O, S, SO_2 or CH_2 .

The β -D- and β -L-nucleosides of this invention may inhibit HCV polymerase activity. Nucleosides can be screened for their ability to inhibit HCV polymerase activity *in vitro* according to screening methods set forth more particularly herein. One can readily determine the spectrum of activity by evaluating the compound in the assays described herein or with another confirmatory assay.

In one embodiment the efficacy of the anti-HCV compound is measured according to the concentration of compound necessary to reduce the plaque number of the virus *in vitro*, according to methods set forth more particularly herein, by 50% (i.e. the compound's

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EC₅₀). In preferred embodiments the compound exhibits an EC₅₀ of less than 25, 15, 10, 5, or 1 micromolar.

In another embodiment, the active compound can be administered in combination or alternation with another anti-HCV agent. In combination therapy, an effective dosage of two or more agents are administered together, whereas during alternation therapy an effective dosage of each agent is administered serially. The dosages will depend on absorption, inactivation, and excretion rates of the drug as well as other factors known to those of skill in the art. It is to be noted that dosage values will also vary with the severity of the condition to be alleviated. It is to be further understood that for any particular subject, specific dosage regimens and schedules should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the compositions.

Nonlimiting examples of antiviral agents that can be used in combination with the compounds disclosed herein include:

(1) an interferon and/or ribavirin (Battaglia, A.M. et al., Ann. Pharmacother. 34:487-494, 2000); Berenguer, M. et al. Antivir. Ther. 3(Suppl. 3):125-136, 1998);

- (2) Substrate-based NS3 protease inhibitors (Attwood et al., Antiviral peptide derivatives, PCT WO 98/22496, 1998; Attwood et al., Antiviral Chemistry and Chemotherapy 10.259-273, 1999; Attwood et al., Preparation and use of amino acid derivatives as anti-viral agents, German Patent Publication DE 19914474; Tung et al. Inhibitors of serine proteases, particularly hepatitis C virus NS3 protease, PCT WO 98/17679), including alphaketoamides and hydrazinoureas, and inhibitors that terminate in an electrophile such as a boronic acid or phosphonate. Llinas-Brunet et al, Hepatitis C inhibitor peptide analogues, PCT WO 99/07734.
- (3) Non-substrate-based inhibitors such as 2,4,6-trihydroxy-3-nitro-benzamide derivatives (Sudo K. et al., Biochemical and Biophysical Research Communications, 238:643-647, 1997; Sudo K. et al. Antiviral Chemistry and Chemotherapy 9:186, 1998), including RD3-4082 and RD3-4078, the former substituted on the amide with a 14 carbon chain and the latter processing a para-phenoxyphenyl group;
- (4) Thiazolidine derivatives which show relevant inhibition in a reverse-phase HPLC assay with an NS3/4A fusion protein and NS5A/5B substrate (Sudo K. et al.,

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Antiviral Research 32:9-18, 1996), especially compound RD-1-6250, possessing a fused cinnamoyl moiety substituted with a long alkyl chain, RD4 6205 and RD4 6193;

- (5) Thiazolidines and benzanilides identified in Kakiuchi N. et al. J. EBS Letters 421:217-220; Takeshita N. et al. Analytical Biochemistry 247:242-246, 1997;
- (6) A phenan-threnequinone possessing activity against HCV protease in a SDS-PAGE and autoradiography assay isolated from the fermentation culture broth of Streptomyces sp., Sch 68631 (Chu M. et al., Tetrahedron Letters 37:7229-7232, 1996), and Sch 351633, isolated from the fungus Penicillium griscofuluum, which demonstrates activity in a scintillation proximity assay (Chu M. et al., Bioorganic and Medicinal Chemistry Letters 9:1949-1952);
- (7) Selective NS3 inhibitors based on the macromolecule elgin c, isolated from leech (Qasim M.A. et al., Biochemistry 36:1598-1607, 1997);
- (8) HCV helicase inhibitors (Diana G.D. et al., Compounds, compositions and methods for treatment of hepatitis C, U.S. Patent No. 5,633,358; Diana G.D. et al., Piperidine derivatives, pharmaceutical compositions thereof and their use in the treatment of hepatitis C, PCT WO 97/36554);
- (9) HCV polymerase inhibitors such as nucleotide analogues, gliotoxin (Ferrari R. et al. Journal of Virology 73:1649-1654, 1999), and the natural product cerulenin (Lohmann V. et al., Virology 249:108-118, 1998);
- (10) Antisense phosphorothioate oligodeoxynucleotides (S-ODN) complementary to sequence stretches in the 5' non-coding region (NCR) of the HCV (Alt M. et al., Hepatology 22:707-717, 1995), or nucleotides 326-348 comprising the 3' end of the NCR and nucleotides 371-388 located in the core coding region of the IICV RNA (Alt M. et al., Archives of Virology 142:589-599, 1997; Galderisi U. et al., Journal of Cellular Physiology 181:251-257, 1999);
- (11) Inhibitors of IRES-dependent translation (Ikeda N et al., Agent for the prevention and treatment of hepatitis C, Japanese Patent Publication JP-08268890; Kai Y. et al. Prevention and treatment of viral diseases, Japanese Patent Publication JP-10101591);

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(12) Nuclease-resistant ribozymes (Maccjak D.J. et al., Hepatology 30 abstract 995, 1999); and

(13) Other miscellaneous compounds including 1-amino-alkylcyclohexanes (U.S. Patent No. 6,034,134 to Gold et al.), alkyl lipids (U.S. Patent No. 5,922,757 to Chojkier et al.), vitamin E and other antioxidants (U.S. Patent No. 5,922,757 to Chojkier et al.), squalene, amantadine, bile acids (U.S. Patent No. 5,846,964 to Ozeki et al.), N-(phosphonoacetyl)-L-aspartic acid, (U.S. Patent No. 5,830,905 to Diana et al.), benzenedicarboxamides (U.S. Patent No. 5,633,388 to Diana et al.), polyadenylic acid derivatives (U.S. Patent No. 5,496,546 to Wang et al.), 2',3'-dideoxyinosine (U.S. Patent No. 5,026,687 to Yarchoan et al.), and benzimidazoles (U.S. Patent No. 5,891,874 to Colacino et al.).

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 provides the structure of various non-limiting examples of nucleosides of the present invention, as well as other known nucleosides, FIAU and Ribavirin, which are used as comparative examples in the text.

Figure 2 is a line graph of the pharmacokinetics (plasma concentrations) of β -D-2'-CH₃-riboG administered to six Cynomolgus Monkeys over time after administration.

Figure 3a and 3b are line graphs of the pharmacokinetics (plasma concentrations) of β -D-2'-CH₃-riboG administered to Cynomolgus Monkeys either intravenously (3a) or orally (3b) over time after administration.

DETAILED DESCRIPTION OF THE INVENTION

The invention as disclosed herein is a compound, method and composition for the treatment of hepatitis C in humans or other host animals, that includes administering an effective HCV treatment amount of a β -D- or β -L-nucleoside as described herein or a pharmaceutically acceptable salt or prodrug thereof, optionally in a pharmaceutically acceptable carrier. The compounds of this invention either possess antiviral (i.e., anti-HCV) activity, or are metabolized to a compound that exhibits such activity.

In summary, the present invention includes the following features:

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(a) β -D- and β -L-nucleosides, as described herein, and pharmaceutically acceptable salts and prodrugs thereof;

- (b) β -D- and β -L-nucleosides as described herein, and pharmaceutically acceptable salts and prodrugs thereof for use in the treatment or prophylaxis of an HCV infection, especially in individuals diagnosed as having an HCV infection or being at risk for becoming infected by HCV;
- (c) use of these β -D- and β -L-nucleosides, and pharmaceutically acceptable salts and prodrugs thereof in the manufacture of a medicament for treatment of an HCV infection;
- (d) pharmaceutical formulations comprising the β -D- or β -L-nucleosides or pharmaceutically acceptable salts or prodrugs thereof together with a pharmaceutically acceptable carrier or diluent;
- (e) β -D- and β -L-nucleosides as described herein substantially in the absence of enantiomers of the described nucleoside, or substantially isolated from other chemical entities;
- (f) processes for the preparation of β -D- and β -L-nucleosides, as described in more detail below; and
- (g) processes for the preparation of β -D- and β -L-nucleosides substantially in the absence of enantiomers of the described nucleoside, or substantially isolated from other chemical entities.

I. Active Compound, and Physiologically Acceptable Salts and Prodrugs Thereof

In a first principal embodiment, a compound of Formula I, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

wherein:

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R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a preferred subembodiment, a compound of Formula I, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

20 R^1 , R^2 and R^3 are independently H or phosphate (preferably H);

X¹ is H;

X² is H or NH₂; and

Y is hydrogen, bromo, chloro, fluoro, iodo, NH2 or OH.

In a second principal embodiment, a compound of Formula II, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

5 wherein:

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 R^1 , R^2 and R^3 are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 or R^3 is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

15 X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a preferred subembodiment, a compound of Formula II, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

R¹, R² and R³ are independently H or phosphate (preferably H);

X¹ is H;

X2 is H or NH2; and

Y is hydrogen, bromo, chloro, fluoro, iodo, NH2 or OH.

In a third principal embodiment, a compound of Formula III, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

$$X^{1}$$
 X^{1}
 X^{2}
 X^{1}
 X^{2}
 X^{2}
 X^{2}
 X^{2}
 X^{2}
 X^{3}
 X^{2}
 X^{3}
 X^{2}
 X^{3}
 X^{2}
 X^{3}
 X^{2}

wherein:

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R¹. R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a preferred subembodiment, a compound of Formula III, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

R¹, R² and R³ are independently H or phosphate (preferably H);

 X^{l} is H;

Y is hydrogen, bromo, chloro, fluoro, iodo, NH2 or OH.

In a fourth principal embodiment, a compound of Formula IV, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

$$X^1$$
 X^1
 X^1

wherein:

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R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a preferred subembodiment, a compound of Formula IV, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

R¹, R² and R³ are independently H or phosphate (preferably H);

X1 is H or CH3; and

Y is hydrogen, bromo, chloro, fluoro, iodo, NH2 or OH.

In a fifth principal embodiment, a compound of Formula V, or a pharmaceutically acceptable salt or produce thereof, is provided:

wherein:

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R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a preferred subembodiment, a compound of Formula V, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

 R^1 , R^2 and R^3 are independently H or phosphate (preferably H);

X1 is H or CH3; and

Y is hydrogen, bromo, chloro, fluoro, iodo, NH2 or OH.

In a sixth principal embodiment, a compound of Formula VI, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

5 wherein:

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R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁵; and R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

In a preferred subembodiment, a compound of Formula VI, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

 R^{1} , R^{2} and R^{3} are independently H or phosphate (preferably H);

X1 is H or CH3; and

Y is hydrogen, bromo, chloro, fluoro, iodo, NH2 or OH.

In a seventh principal embodiment, a compound selected from Formulas VII, VIII and IX, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

5 wherein:

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Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and

X is O, S, SO_2 , or CH_2 .

In a first preferred subembodiment, a compound of Formula VII, VIII or IX, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently hydrogen or phosphate;

R⁶ is alkyl; and

25 X is O, S, SO_2 or CH_2 .

In a second preferred subembodiment, a compound of Formula VII, VIII or IX, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are hydrogens;

R⁶ is alkyl; and

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X is O, S, SO₂ or CH₂.

In a third preferred subembodiment, a compound of Formula VII, VIII or IX, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

 R^1 , R^2 and R^3 are independently hydrogen or phosphate;

R⁶ is alkyl; and

X is O.

In a eighth principal embodiment, a compound of Formula X, XI or XII, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other

pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(acyl), -N(acyl), -N(acyl);

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(loweralkyl)₂, -N(acyl)₂; and

X is O, S, SO_2 or CH_2 .

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In a first preferred subembodiment, a compound of Formula X, XI or XII, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently hydrogen or phosphate;

R⁶ is alkyl; and

X is O, S, SO₂ or CH₂.

In a second preferred subembodiment, a compound of Formula X, XI or XII, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are hydrogens;

R⁶ is alkyl; and

X is O, S, SO_2 or CH_2 .

In a third preferred subembodiment, a compound of Formula X, XI or XII, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H or phosphate;

X is O.

In even more preferred subembodiments, a compound of Formula XI, or its pharmaceutically acceptable salt or prodrug, is provided:

wherein:

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Base is a purine or pyrimidine base as defined herein; optionally substituted with an amine or cyclopropyl (e.g., 2-amino, 2,6-diamino or cyclopropyl guanosine); and

 R^1 and R^2 are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 or R^2 is independently H or phosphate.

In a ninth principal embodiment a compound selected from Formula XIII, XIV or XV, or a pharmaceutically acceptable salt or prodrug thereof, is provided:

wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² or R³ is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -NH(acyl), -NH(acyl), -NH(acyl), -N(acyl)₂; and

X is O, S, SO₂ or CH₂.

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In a first preferred subembodiment, a compound of Formula XIII, XIV or XV, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

 R^1 , R^2 and R^3 are independently hydrogen or phosphate;

R⁶ is alkyl; and

X is O, S, SO_2 or CH_2 .

In a second preferred subembodiment, a compound of Formula XIII, XIV or XV, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are hydrogens;

R⁶ is alkyl; and

25 $X \text{ is } O, S, SO_2 \text{ or } CH_2.$

In a third preferred subembodiment, a compound of Formula XIII, XIV or XV, or a pharmaceutically acceptable salt or prodrug thereof, is provided wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently hydrogen or phosphate;

R⁶ is alkyl; and

X is O.

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In a tenth principal embodiment the invention provides a compound of Formula XVI, or a pharmaceutically acceptable salt or prodrug thereof:

$$R^{10}$$
 X R^{8} R^{6} R^{9} R^{7} (XVI)

wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -NH(acyl), -N(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine;

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alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a pi bond; and

X is O, S, SO₂ or CH₂.

In a first preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR², alkyl, alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (5) R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O, S, SO₂ or CH₂.

In a second preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl, alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O, S, SO₂ or CH₂.

In a third preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl, alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR², alkyl, alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (5) R⁸ and R¹⁰ are H; and (6) X is O, S, SO₂ or CH₂.

In a fourth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl, alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR², alkyl, alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O.

In a fifth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including

monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR¹; (5) R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O, S, SO₂ or CH₂.

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In a sixth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R⁸ and R¹⁰ are H; and (6) X is O, S, SO₂, or CH₂.

In a seventh preferred subembodiment, a compound of Formula XVI, or its

pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including

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monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in

vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3)

R⁶ is alkyl: (4) R⁷ and R⁹ are independently OR², alkyl (including lower alkyl), alkenyl,

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alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO_2 , amino, loweralkylamino or di(loweralkyl)amino; (5) R^8 and R^{10} are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O.

In a eighth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O, S, SO₂ or CH₂.

In a ninth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O.

In a tenth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or

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pyrimidine base as defined herein; (2) R¹ is independently H or phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O.

In an eleventh preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O, S, SO₂ or CH₂.

In a twelfth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O, S, SO₂, or CH₂.

In a thirteenth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O.

In a fourteenth preferred subembodiment, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl;

(4) R⁷ and R⁹ are independently OR², alkyl (including lower alkyl), alkenyl, alkynyl, Brvinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O.

In even more preferred subembodiments, a compound of Formula XVI, or its pharmaceutically acceptable salt or prodrug, is provided in which:

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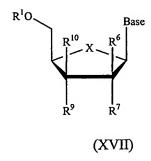
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- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
- (1) Base is guanine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
- (1) Base is cytosine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
 - (1) Base is thymine; (2) R^1 is hydrogen; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^8 and R^{10} are hydrogen; and (6) X is O;
 - (1) Base is uracil; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
 - (1) Base is adenine; (2) R^1 is phosphate; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^8 and R^{10} are hydrogen; and (6) X is O;
 - (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is ethyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is propyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
 - (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is butyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ is hydrogen and R⁹ is hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is O;
 - (1) Base is adenine; (2) R^1 is hydrogen; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^8 and R^{10} are hydrogen; and (6) X is S;
 - (1) Base is adenine; (2) R^1 is hydrogen; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^8 and R^{10} are hydrogen; and (6) X is SO₂;

(1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ and R¹⁰ are hydrogen; and (6) X is CH₂;

In a eleventh principal embodiment the invention provides a compound of Formula XVII, or a pharmaceutically acceptable salt or prodrug thereof:



wherein:

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Base is a purine or pyrimidine base as defined herein;

R¹ is H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

25 R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a pi bond; and

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In a first preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)-amino; (5) R¹⁰ is H; and (6) X is O, S, SO₂, or CH₂.

In a second preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O, S, SO₂ or CH₂.

In a third preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or

pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)-amino; (5) R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O.

In a fourth preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R¹⁰ is H; and (6) X is O, S, SO₂ or CH₂.

In a fifth preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is

optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O.

In a sixth preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R¹⁰ is H; and (6) X is O.

In a seventh preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3)

R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R¹⁰ is H; and (6) X is O.

In an eighth preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)-amino; (5) R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O, S, SO₂, or CH₂.

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In a ninth preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R¹⁰ is H; and (6) X is O, S, SO₂, or CH₂.

In a tenth preferred subembodiment, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR²; (5) R¹⁰ is H; and (6) X is O, S, SO₂, or CH₂.

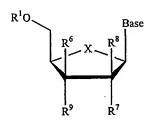
In even more preferred subembodiments, a compound of Formula XVII, or its pharmaceutically acceptable salt or prodrug, is provided in which:

- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;
- (1) Base is guanine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;
- (1) Base is cytosine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;
- (1) Base is thymine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;

(1) Base is uracil; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;

- (1) Base is adenine; (2) R^1 is phosphate; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^{10} is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is ethyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;
 - (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is propyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is butyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is O;
 - (1) Base is adenine; (2) R^1 is hydrogen; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^{10} is hydrogen; and (6) X is S;
 - (1) Base is adenine; (2) R^1 is hydrogen; (3) R^6 is methyl; (4) R^7 and R^9 are hydroxyl; (5) R^{10} is hydrogen; and (6) X is SO₂; or
 - (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R¹⁰ is hydrogen; and (6) X is CH₂.

In an twelfth principal embodiment the invention provides a compound of Formula XVIII, or a pharmaceutically acceptable salt or prodrug thereof:



(XVIII)

wherein:

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Base is a purine or pyrimidine base as defined herein;

R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl,

wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Brvinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(acyl), -N(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, lower alkylamino, or di(loweralkyl)amino;

 R^8 is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R^7 and R^9 , or R^8 and R^9 can come together to form a pi bond;

X is O, S, SO₂ or CH₂.

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In a first preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (5) R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; and (6) X is O, S, SO₂ or CH₂.

In a second preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or

pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di-(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O, S, SO₂ or CH₂.

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In a third preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(lower-alkyl)amino; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R⁸ is H; and (6) X is O, S, SO₂ or CH₂.

In a fourth preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is

optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R⁸ is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O.

In a fifth preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ is H; and (6) X is O, S, SO₂, or CH₂.

In a sixth preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3)

R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; and (6) X is O.

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In a seventh preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered in vivo is capable of providing a compound wherein R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino, or di(loweralkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino; (5) R⁸ is H; and (6) X is O.

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In an eighth preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, hydroxy, O-alkyl, O-alkenyl, chloro, bromo, fluoro, iodo, NO₂, amino, loweralkylamino or di(loweralkyl)amino; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ is H; and (6) X is O, S, SO₂ or CH₂.

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In a ninth preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or pyrimidine base as defined herein; (2) R¹ is independently H or phosphate; (3) R⁶ is alkyl; (4) R⁷ and R⁹ are independently OR²; (5) R⁸ is H; and (6) X is O, S, SO₂, or CH₂.

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In a tenth preferred subembodiment, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which: (1) Base is a purine or

WO 01/90121 PCT/US01/16671 pyrimidine base as defined herein; (2) R^1 is independently H or phosphate; (3) R^6 is alkyl; (4) R^7 and R^9 are independently OR^2 ; (5) R^8 is H; and (6) X is O.

In even more preferred subembodiments, a compound of Formula XVIII, or its pharmaceutically acceptable salt or prodrug, is provided in which:

(1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;

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- (1) Base is guanine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is cytosine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is thymine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is uracil; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is phosphate; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is ethyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is propyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R^1 is hydrogen; (3) R^6 is butyl; (4) R^7 and R^9 are hydroxyl; (5) R^8 is hydrogen; and (6) X is O;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is S;
- (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is SO₂; or
 - (1) Base is adenine; (2) R¹ is hydrogen; (3) R⁶ is methyl; (4) R⁷ and R⁹ are hydroxyl; (5) R⁸ is hydrogen; and (6) X is CH₂.

The β -D- and β -L-nucleosides of this invention may inhibit HCV polymerase activity. Nucleosides can be screened for their ability to inhibit HCV polymerase activity in vitro according to screening methods set forth more particularly herein. One can readily determine the spectrum of activity by evaluating the compound in the assays described herein or with another confirmatory assay.

In one embodiment the efficacy of the anti-HCV compound is measured according to the concentration of compound necessary to reduce the plaque number of the virus *in vitro*, according to methods set forth more particularly herein, by 50% (i.e. the compound's EC₅₀). In preferred embodiments the compound exhibits an EC₅₀ of less than 15 or 10 micromolar, when measured according to the polymerase assay described in Ferrari *et al.*, *Inl. of Vir.*, 73:1649-1654, 1999; Ishii *et al.*, *Hepatology*, 29:1227-1235,1999; Lohmann *et al.*, *Inl. of Bio. Chem.*, 274:10807-10815, 1999; or Yamashita *et al., Inl. of Bio. Chem.*, 273:15479-15486, 1998.

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The active compound can be administered as any salt or prodrug that upon administration to the recipient is capable of providing directly or indirectly the parent compound, or that exhibits activity itself. Nonlimiting examples are the pharmaceutically acceptable salts (alternatively referred to as "physiologically acceptable salts"), and a compound that has been alkylated or acylated at the 5'-position or on the purine or pyrimidine base (a type of "pharmaceutically acceptable prodrug"). Further, the modifications can affect the biological activity of the compound, in some cases increasing the activity over the parent compound. This can easily be assessed by preparing the salt or prodrug and testing its antiviral activity according to the methods described herein, or other methods known to those skilled in the art.

II. Definitions

The term alkyl, as used herein, unless otherwise specified, refers to a saturated straight, branched, or cyclic, primary, secondary, or tertiary hydrocarbon of typically C_1 to C_{10} , and specifically includes methyl, ethyl, propyl, isopropyl, cyclopropyl, butyl, isobutyl, t-butyl, pentyl, cyclopentyl, isopentyl, neopentyl, hexyl, isohexyl, cyclohexyl,

cyclohexylmethyl, 3-methylpentyl, 2,2-dimethylbutyl, and 2,3-dimethylbutyl. The term includes both substituted and unsubstituted alkyl groups. Moieties with which the alkyl group can be substituted are selected from the group consisting of hydroxyl, amino, alkylamino, arylamino, alkoxy, aryloxy, nitro, cyano, sulfonic acid, sulfate, phosphonic acid, phosphate, or phosphonate, either unprotected, or protected as necessary, as known to those skilled in the art, for example, as taught in Greene, et al., Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, hereby incorporated by reference.

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The term lower alkyl, as used herein, and unless otherwise specified, refers to a C₁ to C₄ saturated straight, branched, or if appropriate, a cyclic (for example, cyclopropyl) alkyl group, including both substituted and unsubstituted forms. Unless otherwise specifically stated in this application, when alkyl is a suitable moiety, lower alkyl is preferred. Similarly, when alkyl or lower alkyl is a suitable moiety, unsubstituted alkyl or lower alkyl is preferred.

The term alkylamino or arylamino refers to an amino group that has one or two alkyl or aryl substituents, respectively.

The term "protected" as used herein and unless otherwise defined refers to a group that is added to an oxygen, nitrogen, or phosphorus atom to prevent its further reaction or for other purposes. A wide variety of oxygen and nitrogen protecting groups are known to those skilled in the art of organic synthesis.

The term aryl, as used herein, and unless otherwise specified, refers to phenyl, biphenyl, or naphthyl, and preferably phenyl. The term includes both substituted and unsubstituted moieties. The aryl group can be substituted with one or more moieties selected from the group consisting of hydroxyl, amino, alkylamino, arylamino, alkoxy, aryloxy, nitro, cyano, sulfonic acid, sulfate, phosphonic acid, phosphate, or phosphonate, either unprotected, or protected as necessary, as known to those skilled in the art, for example, as taught in Greene, et al., Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

The term alkaryl or alkylaryl refers to an alkyl group with an aryl substituent. The term aralkyl or arylalkyl refers to an aryl group with an alkyl substituent.

The term halo, as used herein, includes chloro, bromo, iodo, and fluoro.

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The term purine or pyrimidine base includes, but is not limited to, adenine, N⁶alkylpurines, N⁶-acylpurines (wherein acyl is C(O)(alkyl, aryl, alkylaryl, or arylalkyl), N⁶benzylpurine, N⁶-halopurine, N⁶-vinylpurine, N⁶-acetylenic purine, N⁶-acyl purine, N⁶-hydroxyalkyl purine, N⁶-thioalkyl purine, N²-alkylpurines, N²-alkyl-6-thiopurines, thymine, cytosine, 5-fluorocytosine, 5-methylcytosine, 6-azapyrimidine, including 6-azacytosine, 2- and/or 4-mercaptopyrmidine, uracil, 5-halouracil, including 5-fluorouracil, C⁵-alkylpyrimidines, C⁵-benzylpyrimidines, C⁵-halopyrimidines, C⁵-vinylpyrimidine, C⁵acetylenic pyrimidine, C⁵-acyl pyrimidine, C⁵-hydroxyalkyl purine, C⁵-amidopyrimidine, C⁵-cyanopyrimidine, C⁵-nitropyrimidine, C⁵-aminopyrimidine, N²-alkylpurines, N²-alkyl-5-azacytidinyl, 5-azauracilyl, triazolopyridinyl, imidazolopyridinyl, 6-thiopurines, pyrrolopyrimidinyl, and pyrazolopyrimidinyl. Purine bases include, but are not limited to, guanine, adenine, hypoxanthine, 2,6-diaminopurine, and 6-chloropurine. Functional oxygen and nitrogen groups on the base can be protected as necessary or desired. Suitable protecting groups are well known to those skilled in the art, and include trimethylsilyl, dimethylhexylsilyl, t-butyldimethylsilyl, and t-butyldiphenylsilyl, trityl, alkyl groups, and acyl groups such as acetyl and propionyl, methanesulfonyl, and p-toluenesulfonyl.

The term acyl refers to a carboxylic acid ester in which the non-carbonyl moiety of the ester group is selected from straight, branched, or cyclic alkyl or lower alkyl, alkoxyalkyl including methoxymethyl, aralkyl including benzyl, aryloxyalkyl such as phenoxymethyl, aryl including phenyl optionally substituted with chloro, bromo, fluoro, iodo, C₁ to C₄ alkyl or C₁ to C₄ alkoxy, sulfonate esters such as alkyl or aralkyl sulphonyl including methanesulfonyl, the mono, di or triphosphate ester, trityl or monomethoxytrityl, substituted benzyl, trialkylsilyl (e.g. dimethyl-t-butylsilyl) or diphenylmethylsilyl. Aryl groups in the esters optimally comprise a phenyl group. The term "lower acyl" refers to an acyl group in which the non-carbonyl moiety is a lower alkyl.

As used herein, the term "substantially free of" or "substantially in the absence of" refers to a nucleoside composition that includes at least 85 or 90% by weight, preferably 95% to 98 % by weight, and even more preferably 99% to 100% by weight, of the designated enantiomer of that nucleoside. In a preferred embodiment, in the methods and compounds of this invention, the compounds are substantially free of enantiomers.

Similarly, the term "isolated" refers to a nucleoside composition that includes at least 85 or 90% by weight, preferably 95% to 98 % by weight, and even more preferably 99% to 100% by weight, of the nucleoside, the remainder comprising other chemical species or enantiomers.

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The term "independently" is used herein to indicate that the variable which is independently applied varies independently from application to application. Thus, in a compound such as R"XYR", wherein R" is "independently carbon or nitrogen," both R" can be carbon, both R" can be nitrogen, or one R" can be carbon and the other R" nitrogen.

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The term host, as used herein, refers to an unicellular or multicellular organism in which the virus can replicate, including cell lines and animals, and preferably a human. Alternatively, the host can be carrying a part of the hepatitis C viral genome, whose replication or function can be altered by the compounds of the present invention. The term host specifically refers to infected cells, cells transfected with all or part of the HCV genome and animals, in particular, primates (including chimpanzees) and humans. In most animal applications of the present invention, the host is a human patient. Veterinary applications, in certain indications, however, are clearly anticipated by the present invention (such as chimpanzees).

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The term "pharmaceutically acceptable salt or prodrug" is used throughout the specification to describe any pharmaceutically acceptable form (such as an ester, phosphate ester, salt of an ester or a related group) of a nucleoside compound which, upon administration to a patient, provides the nucleoside compound. Pharmaceutically acceptable salts include those derived from pharmaceutically acceptable inorganic or organic bases and acids. Suitable salts include those derived from alkali metals such as potassium and sodium, alkaline earth metals such as calcium and magnesium, among numerous other acids well known in the pharmaceutical art. Pharmaceutically acceptable prodrugs refer to a compound that is metabolized, for example hydrolyzed or oxidized, in the host to form the compound of the present invention. Typical examples of prodrugs include compounds that have biologically labile protecting groups on a functional moiety of the active compound. Prodrugs include compounds that can be oxidized, reduced, aminated, deaminated, hydroxylated, dehydroxylated, hydrolyzed, dehydrolyzed, alkylated, dealkylated, acylated, deacylated, phosphorylated, dephosphorylated to produce the active

compound. The compounds of this invention possess antiviral activity against HCV, or are metabolized to a compound that exhibits such activity.

III. Nucleotide Salt or Prodrug Formulations

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In cases where compounds are sufficiently basic or acidic to form stable nontoxic acid or base salts, administration of the compound as a pharmaceutically acceptable salt may be appropriate. Examples of pharmaceutically acceptable salts are organic acid addition salts formed with acids, which form a physiological acceptable anion, for example, tosylate, methanesulfonate, acetate, citrate, malonate, tartarate, succinate, benzoate, ascorbate, α -ketoglutarate, and α -glycerophosphate. Suitable inorganic salts may also be formed, including, sulfate, nitrate, bicarbonate, and carbonate salts.

Pharmaceutically acceptable salts may be obtained using standard procedures well known in the art, for example by reacting a sufficiently basic compound such as an amine with a suitable acid affording a physiologically acceptable anion. Alkali metal (for example, sodium, potassium or lithium) or alkaline earth metal (for example calcium) salts of carboxylic acids can also be made.

Any of the nucleosides described herein can be administered as a nucleotide prodrug to increase the activity, bioavailability, stability or otherwise alter the properties of the nucleoside. A number of nucleotide prodrug ligands are known. In general, alkylation, acylation or other lipophilic modification of the mono, di or triphosphate of the nucleoside will increase the stability of the nucleotide. Examples of substituent groups that can replace one or more hydrogens on the phosphate moiety are alkyl, aryl, steroids, carbohydrates, including sugars, 1,2-diacylglycerol and alcohols. Many are described in R. Jones and N. Bischofberger, *Antiviral Research*, 27 (1995) 1-17. Any of these can be used in combination with the disclosed nucleosides to achieve a desired effect.

The active nucleoside can also be provided as a 5'-phosphoether lipid or a 5'-ether lipid, as disclosed in the following references, which are incorporated by reference herein: Kucera, L.S., N. Iyer, E. Leake, A. Raben, Modest E.K., D.L.W., and C. Piantadosi. 1990. "Novel membrane-interactive ether lipid analogs that inhibit infectious HIV-1 production and induce defective virus formation." *AIDS Res. Hum. Retro Viruses*. 6:491-501; Piantadosi, C., J. Marasco C.J., S.L. Morris-Natschke, K.L. Meyer, F. Gumus, J.R. Surles,

K.S. Ishaq, L.S. Kucera, N. Iyer, C.A. Wallen, S. Piantadosi, and E.J. Modest. 1991. "Synthesis and evaluation of novel ether lipid nucleoside conjugates for anti-HIV activity." *J. Med. Chem.* 34:1408.1414; Hosteller, K.Y., D.D. Richman, D.A. Carson, L.M. Stuhmiller, G.M. T. van Wijk, and H. van den Bosch. 1992. "Greatly enhanced inhibition of human immunodeficiency virus type 1 replication in CEM and HT4-6C cells by 3'-deoxythymidine diphosphate dimyristoylglycerol, a lipid prodrug of 3,-deoxythymidine." *Antimicrob. Agents Chemother.* 36:2025.2029; Hosetler, K.Y., L.M. Stuhmiller, H.B. Lenting, H. van den Bosch, and D.D. Richman, 1990. "Synthesis and antiretroviral activity of phospholipid analogs of azidothymidine and other antiviral nucleosides." *J. Biol. Chem.* 265:61127.

Nonlimiting examples of U.S. patents that disclose suitable lipophilic substituents that can be covalently incorporated into the nucleoside, preferably at the 5'-OH position of the nucleoside or lipophilic preparations, include U.S. Patent Nos. 5,149,794 (Sep. 22, 1992, Yatvin et al.); 5,194,654 (Mar. 16, 1993, Hostetler et al., 5,223,263 (June 29, 1993, Hostetler et al.); 5,256,641 (Oct. 26, 1993, Yatvin et al.); 5,411,947 (May 2, 1995, Hostetler et al.); 5,463,092 (Oct. 31, 1995, Hostetler et al.); 5,543,389 (Aug. 6, 1996, Yatvin et al.); 5,543,390 (Aug. 6, 1996, Yatvin et al.); 3,543,391 (Aug. 6, 1996, Yatvin et al.); and 5,554,728 (Sep. 10, 1996; Basava et al.), all of which are incorporated herein by reference. Foreign patent applications that disclose lipophilic substituents that can be attached to the nucleosides of the present invention, or lipophilic preparations, include WO 89/02733, WO 90/00555, WO 91/16920, WO 91/18914, WO 93/00910, WO 94/26273, WO 96/15132, EP 0 350 287, EP 93917054.4, and WO 91/19721.

IV. Combination and Alternation Therapy

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It has been recognized that drug-resistant variants of HCV can emerge after prolonged treatment with an antiviral agent. Drug resistance most typically occurs by mutation of a gene that encodes for an enzyme used in viral replication. The efficacy of a drug against HCV infection can be prolonged, augmented, or restored by administering the compound in combination or alternation with a second, and perhaps third, antiviral compound that induces a different mutation from that caused by the principle drug. Alternatively, the pharmacokinetics, biodistribution or other parameter of the drug can be altered by such combination or alternation therapy. In general, combination therapy is

typically preferred over alternation therapy because it induces multiple simultaneous stresses on the virus.

Nonlimiting examples of antiviral agents that can be used in combination with the compounds disclosed herein include:

(1) an interferon and/or ribavirin (Battaglia, A.M. et al., Ann. Pharmacother. 34:487-494, 2000); Berenguer, M. et al. Antivir. Ther. 3(Suppl. 3):125-136, 1998);

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- (2) Substrate-based NS3 protease inhibitors (Attwood et al., Antiviral peptide derivatives, PCT WO 98/22496, 1998; Attwood et al., Antiviral Chemistry and Chemotherapy 10.259-273, 1999; Attwood et al., Preparation and use of amino acid derivatives as anti-viral agents, German Patent Publication DE 19914474; Tung et al. Inhibitors of serine proteases, particularly hepatitis C virus NS3 protease, PCT WO 98/17679), including alphaketoamides and hydrazinoureas, and inhibitors that terminate in an electrophile such as a boronic acid or phosphonate. Llinas-Brunet et al, Hepatitis C inhibitor peptide analogues, PCT WO 99/07734.
- (3) Non-substrate-based inhibitors such as 2,4,6-trihydroxy-3-nitro-benzamide derivatives (Sudo K. et al., Biochemical and Biophysical Research Communications, 238:643-647, 1997; Sudo K. et al. Antiviral Chemistry and Chemotherapy 9:186, 1998), including RD3-4082 and RD3-4078, the former substituted on the amide with a 14 carbon chain and the latter processing a para-phenoxyphenyl group;
- (4) Thiazolidine derivatives which show relevant inhibition in a reverse-phase HPLC assay with an NS3/4A fusion protein and NS5A/5B substrate (Sudo K. et al., Antiviral Research 32:9-18, 1996), especially compound RD-1-6250, possessing a fused cinnamoyl moiety substituted with a long alkyl chain, RD4 6205 and RD4 6193;
- (5) Thiazolidines and benzanilides identified in Kakiuchi N. et al. J. EBS Letters 421:217-220; Takeshita N. et al. Analytical Biochemistry 247:242-246, 1997;
- (6) A phenan-threnequinone possessing activity against HCV protease in a SDS-PAGE and autoradiography assay isolated from the fermentation culture broth of Streptomyces sp., Sch 68631 (Chu M. et al., Tetrahedron Letters 37:7229-7232, 1996), and Sch 351633, isolated from the fungus Penicillium griscofuluum, which demonstrates activity in a scintillation proximity assay (Chu M. et al., Bioorganic and Medicinal Chemistry Letters 9:1949-1952);

(7) Selective NS3 inhibitors based on the macromolecule elgin c, isolated from leech (Qasim M.A. et al., Biochemistry 36:1598-1607, 1997);

(8) HCV helicase inhibitors (Diana G.D. et al., Compounds, compositions and methods for treatment of hepatitis C, U.S. Patent No. 5,633,358; Diana G.D. et al., Piperidine derivatives, pharmaceutical compositions thereof and their use in the treatment of hepatitis C, PCT WO 97/36554);

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- (9) HCV polymerase inhibitors such as nucleotide analogues, gliotoxin (Ferrari R. et al. Journal of Virology 73:1649-1654, 1999), and the natural product cerulenin (Lohmann V. et al., Virology 249:108-118, 1998);
- (10) Antisense phosphorothioate oligodeoxynucleotides (S-ODN) complementary to sequence stretches in the 5' non-coding region (NCR) of the HCV (Alt M. et al., Hepatology 22:707-717, 1995), or nucleotides 326-348 comprising the 3' end of the NCR and nucleotides 371-388 located in the core coding region of the IICV RNA (Alt M. et al., Archives of Virology 142:589-599, 1997; Galderisi U. et al., Journal of Cellular Physiology 181:251-257, 1999);
- (11) Inhibitors of IRES-dependent translation (Ikeda N et al., Agent for the prevention and treatment of hepatitis C, Japanese Patent Publication JP-08268890; Kai Y. et al. Prevention and treatment of viral diseases, Japanese Patent Publication JP-10101591);
- (12) Nuclease-resistant ribozymes. (Maccjak D.J. et al., Hepatology 30 abstract 995, 1999); and
- (13) Other miscellaneous compounds including 1-amino-alkylcyclohexanes (U.S. Patent No. 6,034,134 to Gold *et al.*), alkyl lipids (U.S. Patent No. 5,922,757 to Chojkier *et al.*), vitamin E and other antioxidants (U.S. Patent No. 5,922,757 to Chojkier *et al.*), squalene, amantadine, bile acids (U.S. Patent No. 5,846,964 to Ozeki *et al.*), N-(phosphonoacetyl)-L-aspartic acid, (U.S. Patent No. 5,830,905 to Diana *et al.*), benzenedicarboxamides (U.S. Patent No. 5,633,388 to Diana *et al.*), polyadenylic acid derivatives (U.S. Patent No. 5,496,546 to Wang *et al.*), 2',3'-dideoxyinosine (U.S. Patent No. 5,026,687 to Yarchoan *et al.*), and benzimidazoles (U.S. Patent No. 5,891,874 to Colacino *et al.*).

V. Pharmaceutical Compositions

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Hosts, including humans, infected with HCV, or a gene fragment thereof, can be treated by administering to the patient an effective amount of the active compound or a pharmaceutically acceptable prodrug or salt thereof in the presence of a pharmaceutically acceptable carrier or diluent. The active materials can be administered by any appropriate route, for example, orally, parenterally, intravenously, intradermally, subcutaneously, or topically, in liquid or solid form.

A preferred dose of the compound for HCV will be in the range from about 1 to 50 mg/kg, preferably 1 to 20 mg/kg, of body weight per day, more generally 0.1 to about 100 mg per kilogram body weight of the recipient per day. The effective dosage range of the pharmaceutically acceptable salts and prodrugs can be calculated based on the weight of the parent nucleoside to be delivered. If the salt or prodrug exhibits activity in itself, the effective dosage can be estimated as above using the weight of the salt or prodrug, or by other means known to those skilled in the art.

The compound is conveniently administered in unit any suitable dosage form, including but not limited to one containing 7 to 3000 mg, preferably 70 to 1400 mg of active ingredient per unit dosage form. A oral dosage of 50-1000 mg is usually convenient.

Ideally the active ingredient should be administered to achieve peak plasma concentrations of the active compound of from about 0.2 to 70 μ M, preferably about 1.0 to 10 μ M. This may be achieved, for example, by the intravenous injection of a 0.1 to 5% solution of the active ingredient, optionally in saline, or administered as a bolus of the active ingredient.

The concentration of active compound in the drug composition will depend on absorption, inactivation and excretion rates of the drug as well as other factors known to those of skill in the art. It is to be noted that dosage values will also vary with the severity of the condition to be alleviated. It is to be further understood that for any particular subject, specific dosage regimens should be adjusted over time according to the individual need and the professional judgment of the person administering or supervising the administration of the compositions, and that the concentration ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the claimed composition. The active ingredient may be administered at once, or may be divided into a number of smaller doses to be administered at varying intervals of time.

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A preferred mode of administration of the active compound is oral. Oral compositions will generally include an inert diluent or an edible carrier. They may be enclosed in gelatin capsules or compressed into tablets. For the purpose of oral therapeutic administration, the active compound can be incorporated with excipients and used in the form of tablets, troches, or capsules. Pharmaceutically compatible binding agents, and/or adjuvant materials can be included as part of the composition.

The tablets, pills, capsules, troches and the like can contain any of the following ingredients, or compounds of a similar nature: a binder such as microcrystalline cellulose, gum tragacanth or gelatin; an excipient such as starch or lactose, a disintegrating agent such as alginic acid, Primogel, or corn starch; a lubricant such as magnesium stearate or Sterotes; a glidant such as colloidal silicon dioxide; a sweetening agent such as sucrose or saccharin; or a flavoring agent such as peppermint, methyl salicylate, or orange flavoring. When the dosage unit form is a capsule, it can contain, in addition to material of the above type, a liquid carrier such as a fatty oil. In addition, dosage unit forms can contain various other materials which modify the physical form of the dosage unit, for example, coatings of sugar, shellac, or other enteric agents.

The compound can be administered as a component of an elixir, suspension, syrup, wafer, chewing gum or the like. A syrup may contain, in addition to the active compounds, sucrose as a sweetening agent and certain preservatives, dyes and colorings and flavors.

The compound or a pharmaceutically acceptable prodrug or salts thereof can also be mixed with other active materials that do not impair the desired action, or with materials that supplement the desired action, such as antibiotics, antifungals, anti-inflammatories, or other antivirals, including other nucleoside compounds. Solutions or suspensions used for parenteral, intradermal, subcutaneous, or topical application can include the following components: a sterile diluent such as water for injection, saline solution, fixed oils, polyethylene glycols, glycerine, propylene glycol or other synthetic solvents; antibacterial agents such as benzyl alcohol or methyl parabens; antioxidants such as ascorbic acid or sodium bisulfite; chelating agents such as ethylenediaminetetraacetic acid; buffers such as acetates, citrates or phosphates and agents for the adjustment of tonicity such as sodium chloride or dextrose. The parental preparation can be enclosed in ampoules, disposable syringes or multiple dose vials made of glass or plastic.

If administered intravenously, preferred carriers are physiological saline or phosphate buffered saline (PBS).

In a preferred embodiment, the active compounds are prepared with carriers that will protect the compound against rapid elimination from the body, such as a controlled release formulation, including implants and microencapsulated delivery systems. Biodegradable, biocompatible polymers can be used, such as ethylene vinyl acetate, polyanhydrides, polyglycolic acid, collagen, polyorthoesters and polylactic acid. Methods for preparation of such formulations will be apparent to those skilled in the art. The materials can also be obtained commercially from Alza Corporation.

Liposomal suspensions (including liposomes targeted to infected cells with monoclonal antibodies to viral antigens) are also preferred as pharmaceutically acceptable carriers. These may be prepared according to methods known to those skilled in the art, for example, as described in U.S. Patent No. 4,522,811 (which is incorporated herein by reference in its entirety). For example, liposome formulations may be prepared by dissolving appropriate lipid(s) (such as stearoyl phosphatidyl ethanolamine, stearoyl phosphatidyl choline, arachadoyl phosphatidyl choline, and cholesterol) in an inorganic solvent that is then evaporated, leaving behind a thin film of dried lipid on the surface of the container. An aqueous solution of the active compound or its monophosphate, diphosphate, and/or triphosphate derivatives is then introduced into the container. The container is then swirled by hand to free lipid material from the sides of the container and to disperse lipid aggregates, thereby forming the liposomal suspension.

VI. Processes for the Preparation of Active Compounds

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The nucleosides of the present invention can be synthesized by any means known in the art. In particular, the synthesis of the present nucleosides can be achieved by either alkylating the appropriately modified sugar, followed by glycosylation or glycosylation followed by alkylation of the nucleoside. The following non-limiting embodiments illustrate some general methodology to obtain the nucleosides of the present invention.

A. General Synthesis of 1'-C-Branched Nucleosides

1'-C-Branched ribonucleosides of the following structure:

wherein BASE is a purine or pyrimidine base as defined herein;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a pi bond;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ or R² is independently H or phosphate;

R⁶ is an alkyl, chloro-, bromo-, fluoro-, or iodo-alkyl (i.e. CF₃), alkenyl, or alkynyl (i.e. allyl); and

X is O, S, SO_2 or CH_2

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can be prepared by one of the following general methods.

1) Modification from the lactone

The key starting material for this process is an appropriately substituted lactone. The lactone can be purchased or can be prepared by any known means including standard epimerization, substitution and cyclization techniques. The lactone can be optionally

protected with a suitable protecting group, preferably with an acyl or silyl group, by methods well known to those skilled in the art, as taught by Greene et al. Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991. The protected lactone can then be coupled with a suitable coupling agent, such as an organometallic carbon nucleophile, such as a Grignard reagent, an organolithium, lithium dialkylcopper or R⁶-SiMe₃ in TBAF with the appropriate non-protic solvent at a suitable temperature, to give the 1'-alkylated sugar.

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The optionally activated sugar can then be coupled to the BASE by methods well known to those skilled in the art, as taught by Townsend Chemistry of Nucleosides and Nucleotides, Plenum Press, 1994. For example, an acylated sugar can be coupled to a silylated base with a lewis acid, such as tin tetrachloride, titanium tetrachloride or trimethylsilyltriflate in the appropriate solvent at a suitable temperature.

Subsequently, the nucleoside can be deprotected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

In a particular embodiment, the 1'-C-branched ribonucleoside is desired. The synthesis of a ribonucleoside is shown in **Scheme 1**. Alternatively, deoxyribo-nucleoside is desired. To obtain these nucleosides, the formed ribonucleoside can optionally be protected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, and then the 2'-OH can be reduced with a suitable reducing agent. Optionally, the 2'-hydroxyl can be activated to facilitate reduction; i.e. via the Barton reduction.

Scheme 1

2. Alternative method for the preparation of 1'-C-branched nucleosides

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The key starting material for this process is an appropriately substituted hexose. The hexose can be purchased or can be prepared by any known means including standard epimerization, such as alkaline treatment, substitution and coupling techniques. The hexose can be selectively protected to give the appropriate hexa-furanose, as taught by Townsend Chemistry of Nucleosides and Nucleotides, Plenum Press, 1994.

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The 1'-hydroxyl can be optionally activated to a suitable leaving group such as an acyl group or a chloro, bromo, fluoro, iodo via acylation or halogenation, respectively. The optionally activated sugar can then be coupled to the BASE by methods well known to those skilled in the art, as taught by Townsend Chemistry of Nucleosides and Nucleotides, Plenum Press, 1994. For example, an acylated sugar can be coupled to a silylated base with a lewis acid, such as tin tetrachloride, titanium tetrachloride or trimethylsilyltriflate in the appropriate solvent at a suitable temperature. Alternatively, a halo-sugar can be coupled to a silylated base with the presence of trimethylsilyltriflate.

The 1'-CH₂-OH, if protected, can be selectively deprotected by methods well known in the art. The resultant primary hydroxyl can be functionalized to yield various C-branched nucleosides. For example, the primary hydroxyl can be reduced to give the methyl, using a suitable reducing agent. Alternatively, the hydroxyl can be activated prior to reduction to facilitate the reaction; i.e. via the Barton reduction. In an alternate embodiment, the primary hydroxyl can be oxidized to the aldehyde, then coupled with a carbon nucleophile, such as a Grignard reagent, an organolithium, lithium dialkylcopper or R⁶-SiMe₃ in TBAF with the appropriate non-protic solvent at a suitable temperature.

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In a particular embodiment, the 1'-C-branched ribonucleoside is desired. The synthesis of a ribonucleoside is shown in Scheme 2. Alternatively, deoxyribo-nucleoside is desired. To obtain these nucleosides, the formed ribonucleoside can optionally be protected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, and then the 2'-OH can be reduced with a suitable reducing agent. Optionally, the 2'-hydroxyl can be activated to facilitate reduction; i.e. via the Barton reduction.

Scheme 2

In addition, the L-enantiomers corresponding to the compounds of the invention can be prepared following the same general methods (1 or 2), beginning with the corresponding L-sugar or nucleoside L-enantiomer as starting material.

B. General Synthesis of 2'-C-Branched Nucleosides

2'-C-Branched ribonucleosides of the following structure:

wherein BASE is a purine or pyrimidine base as defined herein;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a pi bond;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ or R² is independently H or phosphate;

R⁶ is an alkyl, chloro-, bromo-, fluoro-, iodo-alkyl (i.e. CF₃), alkenyl, or alkynyl (i.e. allyl); and

X is O, S, SO₂ or CH₂

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can be prepared by one of the following general methods.

1. Glycosylation of the nucleobase with an appropriately modified sugar

The key starting material for this process is an appropriately substituted sugar with a 2'-OH and 2'-H, with the appropriate leaving group (LG), for example an acyl group or a

chloro, bromo, fluoro or iodo. The sugar can be purchased or can be prepared by any known means including standard epimerization, substitution, oxidation and reduction techniques. The substituted sugar can then be oxidized with the appropriate oxidizing agent in a compatible solvent at a suitable temperature to yield the 2'-modified sugar. Possible oxidizing agents are Jones reagent (a mixture of chromic acid and sulfuric acid), Collins's reagent (dipyridine Cr(VI) oxide, Corey's reagent (pyridinium chlorochromate), pyridinium dichromate, acid dichromate, potassium permanganate, MnO₂, ruthenium tetroxide, phase transfer catalysts such as chromic acid or permanganate supported on a polymer, Cl₂-pyridine, H₂O₂-ammonium molybdate, NaBrO₂-CAN, NaOCl in HOAc, copper chromite, copper oxide, Raney nickel, palladium acetate, Meerwin-Pondorf-Verley reagent (aluminum t-butoxide with another ketone) and N-bromosuccinimide.

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Then coupling of an organometallic carbon nucleophile, such as a Grignard reagent, an organolithium, lithium dialkylcopper or R⁶-SiMe₃ in TBAF with the ketone with the appropriate non-protic solvent at a suitable temperature, yields the 2'-alkylated sugar. The alkylated sugar can be optionally protected with a suitable protecting group, preferably with an acyl or silyl group, by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

The optionally protected sugar can then be coupled to the BASE by methods well known to those skilled in the art, as taught by Townsend Chemistry of Nucleosides and Nucleotides, Plenum Press, 1994. For example, an acylated sugar can be coupled to a silylated base with a lewis acid, such as tin tetrachloride, titanium tetrachloride or trimethylsilyltriflate in the appropriate solvent at a suitable temperature. Alternatively, a halo-sugar can be coupled to a silylated base with the presence of trimethylsilyltriflate.

Subsequently, the nucleoside can be deprotected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

In a particular embodiment, the 2'-C-branched ribonucleoside is desired. The synthesis of a ribonucleoside is shown in Scheme 3. Alternatively, deoxyribo-nucleoside is desired. To obtain these nucleosides, the formed ribonucleoside can optionally be protected by methods well known to those skilled in the art, as taught by Greene et al. Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, and then the 2'-

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OH can be reduced with a suitable reducing agent. Optionally, the 2'-hydroxyl can be

activated to facilitate reduction; i.e. via the Barton reduction.

Scheme 3

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2. Modification of a pre-formed nucleoside

The key starting material for this process is an appropriately substituted nucleoside with a 2'-OH and 2'-H. The nucleoside can be purchased or can be prepared by any known means including standard coupling techniques. The nucleoside can be optionally protected with suitable protecting groups, preferably with acyl or silyl groups, by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

The appropriately protected nucleoside can then be oxidized with the appropriate oxidizing agent in a compatible solvent at a suitable temperature to yield the 2'-modified sugar. Possible oxidizing agents are Jones reagent (a mixture of chromic acid and sulfuric acid), Collins's reagent (dipyridine Cr(VI) oxide, Corey's reagent (pyridinium chlorochromate), pyridinium dichromate, acid dichromate, potassium permanganate, MnO₂,

ruthenium tetroxide, phase transfer catalysts such as chromic acid or permanganate supported on a polymer, Cl₂-pyridine, H₂O₂-ammonium molybdate, NaBrO₂-CAN, NaOCl in HOAc, copper chromite, copper oxide, Raney nickel, palladium acetate, Meerwin-Pondorf-Verley reagent (aluminum *t*-butoxide with another ketone) and *N*-bromosuccinimide.

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Subsequently, the nucleoside can be deprotected by methods well known to those skilled in the art, as taught by GreeneGreene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

In a particular embodiment, the 2'-C-branched ribonucleoside is desired. The synthesis of a ribonucleoside is shown in **Scheme 4**. Alternatively, deoxyribo-nucleoside is desired. To obtain these nucleosides, the formed ribonucleoside can optionally be protected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, and then the 2'-OH can be reduced with a suitable reducing agent. Optionally, the 2'-hydroxyl can be activated to facilitate reduction; i.e. via the Barton reduction.

Scheme 4

In another embodiment of the invention, the L-enantiomers are desired. Therefore, the L-enantiomers can be corresponding to the compounds of the invention can be prepared following the same foregoing general methods, beginning with the corresponding L-sugar or nucleoside L-enantiomer as starting material.

C. General Synthesis of 3'-C-Branched Nucleosides

3'-C-Branched ribonucleosides of the following structure:

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wherein BASE is a purine or pyrimidine base as defined herein;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower

acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a pi bond;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ or R² is independently H or phosphate;

R⁶ is an alkyl, chloro-, fluoro-, bromo-, iodo-alkyl (i.e. CF₃), alkenyl, or alkynyl (i.e. allyl); and

15 X is O, S, SO₂ or CH₂

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can be prepared by one of the following general methods.

1. Glycosylation of the nucleobase with an appropriately modified sugar

The key starting material for this process is an appropriately substituted sugar with a 3'-OH and 3'-H, with the appropriate leaving group (LG), for example an acyl group or a chloro, bromo, fluoro, iodo. The sugar can be purchased or can be prepared by any known means including standard epimerization, substitution, oxidation and reduction techniques. The substituted sugar can then be oxidized with the appropriate oxidizing agent in a compatible solvent at a suitable temperature to yield the 3'-modified sugar. Possible oxidizing agents are Jones reagent (a mixture of chromic acid and sulfuric acid), Collins's reagent (dipyridine Cr(VI) oxide, Corey's reagent (pyridinium chlorochromate), pyridinium dichromate, acid dichromate, potassium permanganate, MnO₂, ruthenium tetroxide, phase transfer catalysts such as chromic acid or permanganate supported on a polymer, Cl₂-

wo 01/90121 PCT/US01/16671 pyridine, H₂O₂-ammonium molybdate, NaBrO₂-CAN, NaOCl in HOAc, copper chromite, copper oxide, Raney nickel, palladium acetate, Meerwin-Pondorf-Verley reagent (aluminum *t*-butoxide with another ketone) and *N*-bromosuccinimide.

Then coupling of an organometallic carbon nucleophile, such as a Grignard reagent, an organolithium, lithium dialkylcopper or R⁶-SiMe₃ in TBAF with the ketone with the appropriate non-protic solvent at a suitable temperature, yields the 3'-C-branched sugar. The 3'-C-branched sugar can be optionally protected with a suitable protecting group, preferably with an acyl or silyl group, by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

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The optionally protected sugar can then be coupled to the BASE by methods well known to those skilled in the art, as taught by Townsend Chemistry of Nucleosides and Nucleotides, Plenum Press, 1994. For example, an acylated sugar can be coupled to a silylated base with a lewis acid, such as tin tetrachloride, titanium tetrachloride or trimethylsilyltriflate in the appropriate solvent at a suitable temperature. Alternatively, a halo-sugar can be coupled to a silylated base with the presence of trimethylsilyltriflate.

Subsequently, the nucleoside can be deprotected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

In a particular embodiment, the 3'-C-branched ribonucleoside is desired. The synthesis of a ribonucleoside is shown in Scheme 5. Alternatively, deoxyribonucleoside is desired. To obtain these nucleosides, the formed ribonucleoside can optionally be protected by methods well known to those skilled in the art, as taught by Greene et al. Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, and then the 2'-OH can be reduced with a suitable reducing agent. Optionally, the 2'-hydroxyl can be activated to facilitate reduction; i.e. via the Barton reduction.

Scheme 5

2. Modification of a pre-formed nucleoside

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The key starting material for this process is an appropriately substituted nucleoside with a 3'-OH and 3'-H. The nucleoside can be purchased or can be prepared by any known means including standard coupling techniques. The nucleoside can be optionally protected with suitable protecting groups, preferably with acyl or silyl groups, by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

The appropriately protected nucleoside can then be oxidized with the appropriate oxidizing agent in a compatible solvent at a suitable temperature to yield the 2'-modified sugar. Possible oxidizing agents are Jones reagent (a mixture of chromic acid and sulfuric acid), Collins's reagent (dipyridine Cr(VI) oxide, Corey's reagent (pyridinium chlorochromate), pyridinium dichromate, acid dichromate, potassium permanganate, MnO₂, ruthenium tetroxide, phase transfer catalysts such

as chromic acid or permanganate supported on a polymer, Cl₂-pyridine, H₂O₂-ammonium molybdate, NaBrO₂-CAN, NaOCl in HOAc, copper chromite, copper oxide, Raney nickel, palladium acetate, Meerwin-Pondorf-Verley reagent (aluminum *t*-butoxide with another ketone) and *N*-bromosuccinimide.

Subsequently, the nucleoside can be deprotected by methods well known to those skilled in the art, as taught by GreeneGreene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991.

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In a particular embodiment, the 3'-C-branched ribonucleoside is desired. The synthesis of a ribonucleoside is shown in **Scheme 6**. Alternatively, deoxyribonucleoside is desired. To obtain these nucleosides, the formed ribonucleoside can optionally be protected by methods well known to those skilled in the art, as taught by Greene *et al.* Protective Groups in Organic Synthesis, John Wiley and Sons, Second Edition, 1991, and then the 2'-OH can be reduced with a suitable reducing agent. Optionally, the 2'-hydroxyl can be activated to facilitate reduction; i.e. via the Barton reduction.

Scheme 6

In another embodiment of the invention, the L-enantiomers are desired. Therefore, the L-enantiomers can be corresponding to the compounds of the invention can be prepared following the same foregoing general methods, beginning with the corresponding L-sugar or nucleoside L-enantiomer as starting material.

5 Examples

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Example 1: Preparation of 1'-C-methylriboadenine via 6-amino-9-(1-deoxy-β-D-psicofuranosyl)purine

As another alternative method of preparation, the title compound could also be prepared according to a published procedure (J. Farkas, and F. Sorm, "Nucleic acid components and their analogues. XCIV. Synthesis of 6-amino-9-(1-deoxy-β-D-psicofuranosyl)purine", Collect. Czech. Chem. Commun. 1967, 32, 2663-2667. J. Farkas", Collect. Czech. Chem. Commun. 1966, 31, 1535) (Scheme 7).

Scheme 7

In a similar manner, but using the appropriate sugar and pyrimidine or purine bases, the following nucleosides of Formula I are prepared.

wherein:

| R ¹ | R ² | R ³ | X ¹ | X ² | Y |
|----------------|----------------|----------------|----------------|----------------|-----------------|
| Н | H | Н | Н | H | H |
| Н | Н | Н | Н | H | NH ₂ |
| H | Н | Н | Н | H | NH-cyclopropyl |
| Н | Н | Н | H | Н | NH-methyl |
| Н | Н | Н | H | H | NH-ethyl |
| Н | Н | Н | H | H | NH-acetyl |
| Н | Н | Н | H | H | OH |
| Н | Н | Н | Н | H | OMe |
| Н | Н | Н | Н | H | OEt |
| Н | Н | Н | H | H | O-cyclopropyl |
| H | Н | Н | Н | H | O-acetyl |
| Н | Н | Н | Н . | H | SH |
| Н | Н | Н | H | H | SMe |
| Н | Н | Н | Н | H | SEt |
| Н | Н | Н | H | H | S-cyclopropyl |
| Н | Н | Н | H | H | F |
| H | Н | Н | H | H | Cl |
| Н | Н | Н | Н | H | Br |
| Н | Н | Н | H | H | I |
| monophosphate | H | Н | H | H | NH ₂ |
| monophosphate | Н | Н | Н | H | NH-acetyl |
| monophosphate | Н | Н | H | Н | NH-cyclopropyl |

| WO 01/90121 R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | X ² | Y Y |
|-------------------------------|----------------|----------------|----------------|----------------|-----------------|
| monophosphate | H | H | H | H | NH-methyl |
| monophosphate | H | H | Н | H | NH-ethyl |
| monophosphate | Н | H | H | H | OH |
| monophosphate | Н | Н | Н | Н | O-acetyl |
| monophosphate | H | H | Н | Н | OMe |
| monophosphate | Н | H | H | H | OEt |
| monophosphate | Н | H | H | Н | O-cyclopropyl |
| monophosphate | H | H | H | Н | SH |
| monophosphate | H | H | H | Н | SMe |
| monophosphate | H | H | Н | Н | SEt |
| monophosphate | Н | H | H | Н | S-cyclopropyl |
| monophosphate | H | H | Н | Н | F |
| monophosphate | Н | Н | H | H | Cl |
| monophosphate | Н | Н | H | H | Br |
| monophosphate | H | Н | H | Н | I |
| diphosphate | H | Н | Н | Н | NH ₂ |
| diphosphate | Н | H | H | Н | NH-acetyl |
| diphosphate | Н | Н | H | H | NH-cyclopropy |
| diphosphate | Н | H | H | H | NH-methyl |
| diphosphate | Н | H | H | H | NH-ethyl |
| diphosphate | Н | Н | Н | H | ОН |
| diphosphate | H | H | H | H | O-acetyl |
| diphosphate | Н | H | H | H | OMe |
| diphosphate | Н | H | H | H | OEt |
| diphosphate | Н | H | H | H | O-cyclopropyl |
| diphosphate | Н | H | H | H | SH |
| diphosphate | Н | H | Н | H | SMe |
| diphosphate | Н | H | Н | H | SEt |
| diphosphate | Н | H | Н | H | S-cyclopropyl |
| diphosphate | Н | H | H | H | F |
| diphosphate | H | Н | Н | H | Cl |

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|----------------|----------------|----------------|-----|----------------|-----------------|
| \mathbb{R}^1 | R ² | R ³ | X1 | X ² | · Y |
| diphosphate | Н | Н | H · | H | Br |
| diphosphate | Н | Н | H | H | I |
| triphosphate | Н | Н | H | H | NH ₂ |
| triphosphate | Н | Н | H | H | NH-acetyl |
| triphosphate | Н | Н | Н | H | NH-cyclopropyl |
| triphosphate | Н | H | H | H | NH-methyl |
| triphosphate | Н | Н | Н | Н | NH-ethyl |
| triphosphate | Н | H | H | H | OH |
| triphosphate | H | Н | H | H | OMe |
| triphosphate | H | H | H | Н | OEt |
| triphosphate | Н | H | H | H | O-cyclopropyl |
| triphosphate | Н | Н | H | H | O-acetyl |
| triphosphate | Н | Н | H | H | SH |
| triphosphate | Н | H | H | Н | SMe |
| triphosphate | Н | H | H | H | SEt |
| triphosphate | Н | Н | H | H | S-cyclopropyl |
| triphosphate | Н | H | H | Н | F |
| triphosphate | Н | Н | H | H | Cl |
| triphosphate | H | H | Н | Н | Br |
| triphosphate | Н | H | H | H | I |
| monophosphate | monophosphate | monophosphate | Н | H | NH ₂ |
| monophosphate | monophosphate | monophosphate | Н | Н | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | Н | Н | OH |
| monophosphate | monophosphate | monophosphate | Н | Н | F |
| monophosphate | monophosphate | monophosphate | H | H | Cl |
| diphosphate | diphosphate | diphosphate | Н | H | NH ₂ |
| diphosphate | diphosphate | diphosphate | Н | Н | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | H | ОН |
| diphosphate | diphosphate | diphosphate | H | H | F |
| diphosphate | diphosphate | diphosphate | H | H | Cl |
| triphosphate | triphosphate | triphosphate | H | H | NH ₂ |

| R ¹ | R ² | R ³ | X ¹ | X^2 | Y |
|----------------|----------------|----------------|-----------------|-------|-----------------|
| triphosphate | triphosphate | triphosphate | Н | Н | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | H | Н | ОН |
| triphosphate | triphosphate | triphosphate | Н | Н | F |
| triphosphate | triphosphate | triphosphate | H | H | Cl |
| H | Н | Н | F | H | NH ₂ |
| Н | Н | Н | F | Н | NH-cyclopropyl |
| H | H | Н | F | H | ОН |
| H | Н | Н | F | H | F |
| H | H | Н | F | Н | Cl |
| H | H | Н | Cl | Н | NH ₂ |
| H | Н . | H | Cl | Н | NH-cyclopropyl |
| H | H | Н | Cl | Н | OH |
| H | H | Н | Cl | H. | F |
| H | Н | H | Cl | Н | CI |
| H | H | H | Br | Н | NH ₂ |
| H | Н | H | Br | Н | NH-cyclopropyl |
| Н | H | Н | Br | Н | OH |
| Н | H | H | Br | H | F |
| H | Н | Н | Br | Н | Cl |
| H | H | Н | NH ₂ | Н | NH ₂ |
| H | H | Н | NH ₂ | Н | NH-cyclopropyl |
| H | H | H | NH ₂ | Н | OH |
| H | H | Н | NH ₂ | Н | F |
| H | Н | Н | NH ₂ | Н | Cl |
| H | Н | Н | SH | H | NH ₂ |
| H | Н | H | SH | Н | NH-cyclopropyl |
| H | H | Н | SH | H | OH |
| H | Н | H | SH | Н | F |
| H | H . | H | SH | H | Cl |
| acetyl | Н | H | Н | Н | NH ₂ |
| acetyl | H | H | H | H | NH-cyclopropyl |

| WO 01/90121 R ¹ | \mathbb{R}^2 | R ³ | X^1 | X^2 | PCT/US01/16671 Y |
|-------------------------------|----------------|----------------|-------|-------|---------------------|
| | | | | | |
| acetyl | H | H | H | Н | OH |
| acetyl | H | H | H | H | F |
| acetyl | H | H | H | H | Cl |
| acetyl | H | H | F | H | NH ₂ |
| acetyl | H | Н | F | H | NH-cyclopropyl |
| acetyl | H | H | F | H | OH |
| acetyl | H | H | F | H | F |
| acetyl | H | H | F | H | Cl |
| H | acetyl | acetyl | H | H | NH ₂ |
| Н | acetyl | acetyl | H | H | NH-cyclopropyl |
| H | acetyl | acetyl | Н | Н | ОН |
| Н | acetyl | acetyl | H | Н | F |
| Н | acetyl | acetyl | Н | H | Cl |
| acetyl | acetyl | acetyl | Н | Н | NH ₂ |
| acetyl | acetyl | acetyl | Н | Н | NH-cyclopropyl |
| acetyl | acetyl | acetyl | Н | Н | ОН |
| acetyl | acetyl | acetyl | H | H | F |
| acetyl | acetyl | acetyl | H | H | Cl |
| monophosphate | acetyl | acetyl | H | Н | NH ₂ |
| monophosphate | acetyl | acetyl | H | Н | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | Н | Н | OH |
| monophosphate | acetyl | acetyl | Н | H | F |
| monophosphate | acetyl | acetyl | H· | H | Cl |
| diphosphate | acetyl | acetyl | H | H | NH ₂ |
| diphosphate | acetyl | acetyl | H | Н | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | H | H | OH |
| diphosphate | acetyl | acetyl | H | H | F |
| diphosphate | acetyl | acetyl | H | H | Cl |
| triphosphate | acetyl | acetyl | H | H | NH ₂ |
| triphosphate | acetyl | acetyl | Н | H | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | H | H | ОН |

| \mathbb{R}^1 | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | \mathbf{X}^2 | Y |
|----------------|----------------|----------------|----------------|-----------------|-----------------|
| triphosphate | acetyl | acetyl | Н | H | F |
| triphosphate | acetyl | acetyl | H | H | Cl |
| H | H | H | Н | NH ₂ | H |
| Н | H | H | H | NH ₂ | NH ₂ |
| Н | Н | H | Н | NH ₂ | NH-cyclopropyl |
| H | H | H | Н | NH ₂ | NH-methyl |
| Н | Н | H | Н | NH ₂ | NH-ethyl |
| H | Н | H | Н | NH ₂ | NH-acetyl |
| Н | Н | H | Н | NH ₂ | OH |
| Н | Н | H | Н | NH ₂ | OMe |
| Н | Н | H | Н | NH ₂ | OEt |
| Н | H | H | Н | NH ₂ | O-cyclopropyl |
| Н | Н | Н | Н | NH ₂ | O-acetyl |
| Н | H | Н | H | NH ₂ | SH |
| Н | H | Н | Н | NH ₂ | SMe |
| Н | H | Н | Н | NH ₂ | SEt |
| Н | H | Н | Н | NH ₂ | S-cyclopropyl |
| Н | Н | Н | Н | NH ₂ | F |
| Н | H | Н | Н | NH ₂ | Cl |
| Н | H | Н | H | NH ₂ | Br |
| Н | Н | Н | Н | NH ₂ | I |
| monophosphate | Н | H | H | NH ₂ | NH ₂ |
| monophosphate | Н | H | Н | NH ₂ | NH-acetyl |
| monophosphate | Н | Н | Н | NH ₂ | NH-cyclopropyl |
| monophosphate | H | H | H | NH ₂ | NH-methyl |
| monophosphate | H | Н | H | NH ₂ | NH-ethyl |
| monophosphate | Н | H | H | NH ₂ | ОН |
| monophosphate | Н | Н | Н | NH ₂ | O-acetyl |
| monophosphate | Н | Н | H | NH ₂ | OMe |
| monophosphate | H | Н | Н | NH ₂ | OEt |
| monophosphate | Н | H | H | NH ₂ | O-cyclopropyl |

| R¹R²monophosphateHmonophosphateHmonophosphateHmonophosphateHmonophosphateHmonophosphateHmonophosphateHmonophosphateH | H H H H | X¹ H H H H | X ² NH ₂ NH ₂ | Y SH SMe SEt S-cyclopropyl |
|--|------------------|-------------|--|----------------------------|
| monophosphate H | H H H H | H H H | NH ₂ NH ₂ NH ₂ | SMe SEt |
| monophosphate H monophosphate H monophosphate H monophosphate H monophosphate H monophosphate H | H H H | H H H | NH ₂ | SEt |
| monophosphate H monophosphate H monophosphate H monophosphate H monophosphate H | H H H | H H | NH ₂ | |
| monophosphate H monophosphate H monophosphate H monophosphate H | H H | Н | | S-cyclopropyl |
| monophosphate H monophosphate H monophosphate H | Н | | NH ₂ | |
| monophosphate H monophosphate H | | H | 1 | F |
| monophosphate H | ŢŢ | 1.1 | NH ₂ | Cl |
| | H | Н | NH ₂ | Br |
| | Н | Н | NH ₂ | I |
| diphosphate H | Н | H | NH ₂ | NH ₂ |
| diphosphate H | Н | H | NH ₂ | NH-acetyl |
| diphosphate H | Н | H | NH ₂ | NH-cyclopropyl |
| diphosphate H | Н | H | NH ₂ | NH-methyl |
| diphosphate H | Н | H | NH ₂ | NH-ethyl |
| diphosphate H | Н | H | NH ₂ | ОН |
| diphosphate H | Н | Н | NH ₂ | O-acetyl |
| diphosphate H | Н | H | NH ₂ | OMe |
| diphosphate H | Н | . H | NH ₂ | OEt |
| diphosphate H | Н | H | NH ₂ | O-cyclopropyl |
| diphosphate H | Н | H | NH ₂ | SH |
| diphosphate H | H | H | NH ₂ | SMe |
| diphosphate H | Н | H | NH ₂ | SEt |
| diphosphate H | Н | H | NH ₂ | S-cyclopropyl |
| diphosphate H | Н | H | NH ₂ | F |
| diphosphate H | Н | H | NH ₂ | Cl |
| diphosphate H | Н | Н | NH ₂ | Br |
| diphosphate H | Н | Н | NH ₂ | I |
| triphosphate H | Н | H | NH ₂ | NH ₂ |
| triphosphate H | Н | Н | NH ₂ | NH-acetyl |
| triphosphate H | H | Н | NH ₂ | NH-cyclopropyl |
| triphosphate H | Н | H | NH ₂ | NH-methyl |
| triphosphate H | H | H | NH ₂ | NH-ethyl |

WO 01/90121 PCT/US01/16671 R^3 $\overline{\mathbf{X}^{\mathbf{I}}}$ $\overline{\mathbf{R^2}}$ $\overline{X^2}$ R^1 Y H H OH H NH_2 triphosphate H NH_2 OMe triphosphate Η Η NH₂ \overline{H} $\overline{\mathbf{H}}$ OEt triphosphate Н H Η H NH_2 O-cyclopropyl triphosphate Η H NH₂ O-acetyl H triphosphate NH₂ SH Η Η Η triphosphate H $\overline{NH_2}$ SMe H Η triphosphate H NH₂ SEt triphosphate H Η H NH₂ S-cyclopropyl triphosphate Η H Н H H NH_2 F triphosphate Н H \overline{NH}_2 Cl H triphosphate Br Η NH_2 triphosphate Η H H H Ī H NH_2 triphosphate monophosphate H NH_2 NH_2 monophosphate monophosphate H NH_2 NH-cyclopropyl monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate Η NH_2 OH F H NH₂ monophosphate monophosphate monophosphate H NH₂ Cl monophosphate monophosphate monophosphate H NH_2 NH_2 diphosphate diphosphate diphosphate diphosphate \mathbf{H} NH_2 NH-cyclopropyl diphosphate diphosphate H OH NH_2 diphosphate diphosphate diphosphate diphosphate F H NH_2 diphosphate diphosphate Cl H NH₂ diphosphate diphosphate diphosphate triphosphate triphosphate H NH_2 NH_2 triphosphate

H NH_2 triphosphate triphosphate NH-cyclopropyl triphosphate triphosphate \mathbf{H} NH_2 OHtriphosphate triphosphate F H $\overline{\mathrm{NH}_{2}}$ triphosphate triphosphate triphosphate H NH_2 Cl triphosphate triphosphate triphosphate F NH_2 H NH_2 H H F NH-cyclopropyl H H H NH_2 H H $\overline{\mathbf{H}}$ F NH_2 OH

| WO 01/90121 | | | | PC | CT/US01/16671 |
|------------------|----------------|----------------|-----------------|-----------------|-----------------|
| \mathbf{R}^{1} | R ² | R ³ | X ¹ | X ² | Y |
| Н | Н | H | F | NH ₂ | F |
| H | Н | Н | F | NH ₂ | Cl |
| Н | Н | Н | Cl | NH ₂ | NH ₂ |
| H | Н | Н | C1 | NH ₂ | NH-cyclopropyl |
| H | H | Н | Cl | NH ₂ | OH |
| Н | H | Н | Cl | NH ₂ | F |
| Н | H | Н | C1 | NH ₂ | Cl |
| Н | H | Н | Br | NH ₂ | NH ₂ |
| H | H | Н | Br | NH ₂ | NH-cyclopropyl |
| Н | H | Н | Br | NH ₂ | OH |
| Н | H | Н | Br | NH ₂ | F |
| H | H | H | Br | NH ₂ | Cl |
| H | H | Н | NH ₂ | NH ₂ | NH ₂ |
| H | H | Н | NH ₂ | NH ₂ | NH-cyclopropyl |
| H | Н | H | NH ₂ | NH ₂ | ОН |
| Н | H | H | NH ₂ | NH ₂ | F |
| Н | H | Н | NH ₂ | NH ₂ | Cl |
| H | Н | H | SH | NH ₂ | NH ₂ |
| Н | H | Н | SH | NH ₂ | NH-cyclopropyl |
| H | Н | Н | SH | NH ₂ | OH |
| H | Н | Н | SH | NH ₂ | F |
| H | H | Н | SH | NH ₂ | Cl |
| acetyl | H | H | Н | NH ₂ | NH ₂ |
| acetyl | H | Н | Н | NH ₂ | NH-cyclopropyl |
| acetyl | H | Н | Н | NH ₂ | OH |
| acetyl | H | Н | H | NH ₂ | F |
| acetyl | H | H | Н | NH ₂ | Cl |
| acetyl | H | Н | F | NH ₂ | NH ₂ |
| acetyl | H | Н | F | NH ₂ | NH-cyclopropyl |
| acetyl | Н | Н | F | NH ₂ | OH |
| acetyl | H | H | F | NH ₂ | F |

| WO 01/90121 R ¹ | R ² | R ³ | X ¹ | X ² | T/US01/16671 Y |
|-------------------------------|----------------|----------------|----------------|-----------------|-------------------|
| acetyl | Н | H | F | NH ₂ | Cl |
| Н | acetyl | acetyl | H | NH ₂ | NH ₂ |
| Н | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| H | acetyl | acetyl | H | NH ₂ | ОН |
| Н | acetyl | acetyl | Н | NH ₂ | F |
| H | acetyl | acetyl | H | NH ₂ | Cl |
| acetyl | acetyl | acetyl | H | NH ₂ | NH ₂ |
| acetyl | acetyl | acetyl | Н | NH ₂ | NH-cyclopropyl |
| acetyl | acetyl | acetyl | H | NH ₂ | ОН |
| acetyl | acetyl | acetyl | H | NH ₂ | F |
| acetyl | acetyl | acetyl | Н | NH ₂ | Cl |
| monophosphate | acetyl | acetyl | Н | NH ₂ | NH ₂ |
| monophosphate | acetyl | acetyl | Н | NH ₂ | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | Н | NH ₂ | ОН |
| monophosphate | acetyl | acetyl | Н | NH ₂ | F |
| monophosphate | acetyl | acetyl | H | NH ₂ | Cl |
| diphosphate | acetyl | acetyl | Н | NH ₂ | NH ₂ |
| diphosphate | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | H | NH ₂ | ОН |
| diphosphate | acetyl | acetyl | H | NH ₂ | F |
| diphosphate | acetyl | acetyl | Н | NH ₂ | Cl |
| triphosphate | acetyl | acetyl | H | NH ₂ | NH ₂ |
| triphosphate | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | H | NH ₂ | OH |
| triphosphate | acetyl | acetyl | H | NH ₂ | F |
| triphosphate | acetyl | acetyl | H | NH ₂ | Cl |
| H | H | Н | Н | Cl | Н |
| Н . | Н | H | Н | Cl | Н |
| H | Н | H | Н | CI | NH ₂ |
| H | Н | H | H | Cl | NH-cyclopropy |
| H | H | H | H | Cl | NH-methyl |

| | WO 01/90121 PCT/US01/16671 | | | | |
|---------------|----------------------------|----------------|-------------|----------------|-----------------|
| Ri | R ² | R ³ | X^1 | X ² | Y |
| Н | Н | Н | H | Cl | NH-ethyl |
| Н | Н | Н | H | Cl | NH-acetyl |
| H | Н | H | H | Cl | OH |
| Н | Н | Н | H | Cl | OMe |
| Н . | Н | H | H | Cl | OEt |
| H | H | Н | Н | C1 | O-cyclopropyl |
| H | Н | Н | Н | Cl | O-acetyl |
| Н | Н | Н | Н | Cl | SH |
| H | Н | Н | Н | Cl | SMe |
| Н | Н | Н | H | Cl | SEt |
| H | H | Н | H | C1 | S-cyclopropyl |
| monophosphate | H | Н | Н | C1 | NH ₂ |
| monophosphate | H | Н | Н | Cl | NH-acetyl |
| monophosphate | Н | Н | Н | Cl | 'NH-cyclopropyl |
| monophosphate | Н | Н | H | Cl | NH-methyl |
| monophosphate | Н | Н | Н | Cl | NH-ethyl |
| monophosphate | H | Н | Н | Cl | ОН |
| monophosphate | Н | Н | Н | C1 | O-acetyl |
| monophosphate | Н | Н | Н | C1 | OMe |
| monophosphate | Н | Н | H | Cl | OEt |
| monophosphate | H | Н | Н | Cl | O-cyclopropyl |
| monophosphate | H | Н | H | Cl | SH |
| monophosphate | Н | Н | H | Cl | SMe |
| monophosphate | Н | Н | H | Cl | SEt |
| monophosphate | Н | H | Н | Cl | S-cyclopropyl |
| diphosphate | Н | H | H | Cl | NH ₂ |
| diphosphate | Н | H | H | Cl | NH-acetyl |
| diphosphate | Н | Н | Н | Cl | NH-cyclopropyl |
| diphosphate | Н | H | Н | Cl | NH-methyl |
| diphosphate | Н | H | H | Cl | NH-ethyl |
| diphosphate | Н | H | Н | Cl | ОН |
| | <u> </u> | | | 4 | |

WO 01/90121 PCT/US01/16671 $\overline{\mathbf{X}^{\mathbf{I}}}$ $\overline{\mathbf{R}^2}$ $\overline{X^2}$ \mathbb{R}^3 H H Η Cl O-acetyl diphosphate H Cl Η Η OMe diphosphate H Н H Cl OEt diphosphate H Cl diphosphate H Η O-cyclopropyl Cl diphosphate H H H SH H H H Cl SMe diphosphate H C1 H H SEt diphosphate Cl Н H H S-cyclopropyl diphosphate Cl NH_2 $\overline{\mathsf{H}}$ H H triphosphate Cl NH-acetyl triphosphate H Η Η Cl H H $\overline{\mathrm{H}}$ NH-cyclopropyl triphosphate Cl Η H NH-methyl H triphosphate Cl Η H Η NH-ethyl triphosphate Cl triphosphate Η H H OH Cl H OMe H triphosphate H H H Cl **OEt** triphosphate Η CI O-cyclopropyl triphosphate Η H Η Cl H H Η O-acetyl triphosphate Η H H Cl SH triphosphate H Cl SMe triphosphate H Η H Cl SEt H H triphosphate H H Cl triphosphate Η S-cyclopropyl Cl NH₂ monophosphate Η monophosphate monophosphate Cl NH-cyclopropyl monophosphate monophosphate monophosphate Η Cl monophosphate monophosphate H OH monophosphate diphosphate H Cl NH_2 diphosphate diphosphate diphosphate H $\overline{C1}$ NH-cyclopropyl diphosphate diphosphate Η \overline{Cl} OH diphosphate diphosphate diphosphate Cl H NH₂ triphosphate triphosphate triphosphate triphosphate triphosphate Η Cl NH-cyclopropyl triphosphate Cl $\overline{\mathrm{H}}$ OH triphosphate triphosphate triphosphate

| WO 01/90121 | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | X^2 | PCT/US01/16671 Y |
|---------------|----------------|----------------|-----------------|-------|---------------------|
| H | H | H | F | Cl | NH ₂ |
| Н | H | H | F | Cl | NH-cyclopropyl |
| Н | Н | H | F | Cl | OH |
| Н | H | Н | Cl | Cl | NH ₂ |
| Н | H | Н | Cl | Cl | NH-cyclopropyl |
| Н | Н | H | Cl | CI | OH |
| Н | H | Н | Br | Cl | NH ₂ |
| Н | H | Н | Br | Cl | NH-cyclopropyl |
| Н | Н | Н | Br | Cl | ОН |
| Н | Н | H | NH ₂ | Cl | NH ₂ |
| Н . | H | H | NH ₂ | Cl | NH-cyclopropyl |
| Н | H | H | NH ₂ | Cl | ОН |
| Н | H | Н | SH | Cl | NH ₂ |
| H | H | H | SH | Cl | NH-cyclopropyl |
| Н | H | Н | SH | Cl | ОН |
| acetyl | H | Н | H | Cl | NH ₂ |
| acetyl | Н | Н | H | Cl | NH-cyclopropyl |
| acetyl | H | Н | H | Cl | ОН |
| acetyl | H | Н | F | Cl | NH ₂ |
| acetyl | H | H | F | Cl | NH-cyclopropyl |
| acetyl | H | H | F | Cl | OH |
| H | acetyl | acetyl | H | Cl | NH ₂ |
| H | acetyl | acetyl | Н | Cl | NH-cyclopropyl |
| Н | acetyl | acetyl | Н | Cl | OH |
| acetyl | acetyl | acetyl | H | Cl | NH ₂ |
| acetyl | acetyl | acetyl | H | CI | NH-cyclopropyl |
| acetyl | acetyl | acetyl | Н | Cl | ОН |
| monophosphate | acetyl | acetyl | Н | Cl | NH ₂ |
| monophosphate | acetyl | acetyl | H | Cl | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | H | Cl | ОН |
| diphosphate | acetyl | acetyl | H | Cl | NH ₂ |

| \mathbb{R}^1 | R ² | R ³ | X ¹ | X ² | Y |
|----------------|----------------|----------------|----------------|----------------|-----------------|
| diphosphate | acetyl | acetyl | H | Cl | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | H | C1 | OH |
| triphosphate | acetyl | acetyl | H | C1 | NH ₂ |
| triphosphate | acetyl | acetyl | H | C1 | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | Н | Cl | OH |
| H | Н | Н | Н | Cl | NH ₂ |
| H | Н | Н | H | Cl | NH-cyclopropyl |
| H | H | Н | Н | Cl | OH |
| Н | Н | Н | H | Br | NH ₂ |
| H | Н | Н | Н | Br | NH-cyclopropyl |
| H | Н | Н | H | Br | OH |

Alternatively, the following nucleosides of Formula IV are prepared, using the appropriate sugar and pyrimidine or purine bases.

wherein:

5

| R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | Y |
|----------------|----------------|----------------|----------------|-----------------|
| H | H | H | Н | Н |
| Н | H | H | Н | NH ₂ |
| H | Н | H | H | NH-cyclopropyl |
| H | Н | Н | H | NH-methyl |
| Н | Н | H | H | NH-ethyl |
| Н | Н | Н | H | NH-acetyl |

WO 01/90121 PCT/US01/16671 $\overline{\mathbf{X}^{\mathbf{I}}}$ $\overline{R^2}$ R^3 $\overline{\mathbf{Y}}$ $\overline{\mathbf{R}^{\mathbf{I}}}$ Н H Η H OH H H H H OMe H H H H OEt H H O-cyclopropyl H Η H Н Н O-acetyl Н H H Η H SH H H H SMe H H SEt H H Η H H H H S-cyclopropyl NH₂ H monophosphate Η Η H H NH-acetyl Н monophosphate H H H NH-cyclopropyl monophosphate H H NH-methyl H monophosphate Η NH-ethyl H Η monophosphate H OH H H monophosphate H O-acetyl monophosphate Η H Η OMe monophosphate Η H H H $\overline{\mathrm{H}}$ OEt monophosphate H H H O-cyclopropyl monophosphate H H SH Η monophosphate H SMe H H monophosphate H SEt H H monophosphate H H S-cyclopropyl Н monophosphate Н H NH₂ diphosphate Η H H NH-acetyl diphosphate H H diphosphate NH-cyclopropyl H Η H H NH-methyl Н diphosphate H diphosphate H H NH-ethyl H OH $\overline{\mathbf{H}}$ diphosphate H O-acetyl diphosphate H H H H H OMe diphosphate H

| WO 01/90121 | | | | PCT/US01/1667 |
|----------------|----------------|----------------|----|-----------------|
| \mathbb{R}^1 | \mathbb{R}^2 | R ³ | X1 | Y |
| diphosphate | Н | H | H | OEt |
| diphosphate | Н | H | Н | O-cyclopropyl |
| diphosphate | H | Н | H | SH |
| diphosphate | Н | H | H | SMe |
| diphosphate | Н | Н | Н | SEt |
| diphosphate | Н | H | Н | S-cyclopropyl |
| triphosphate | Н | Н | Н | NH ₂ |
| triphosphate | Н | Н | Н | NH-acetyl |
| triphosphate | Н | H . | H | NH-cyclopropyl |
| triphosphate | Н | Н | H | NH-methyl |
| triphosphate | Н | H | Н | NH-ethyl |
| triphosphate | H | Н | H | ОН |
| triphosphate | H | H | Н | OMe |
| triphosphate | Н | H | H | OEt |
| triphosphate | Н | Н | Н | O-cyclopropyl |
| triphosphate | Н | Н | Н | O-acetyl |
| triphosphate | Н | Н | H | SH |
| triphosphate | Н | Н | H | SMe |
| triphosphate | Н | H | H | SEt |
| triphosphate | Н | Н | H | S-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | NH ₂ |
| monophosphate | monophosphate | monophosphate | H | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | ОН |
| diphosphate | diphosphate | diphosphate | H | NH ₂ |
| diphosphate | diphosphate | diphosphate | H | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | OH |
| triphosphate | triphosphate | triphosphate | H | NH ₂ |
| triphosphate | triphosphate | triphosphate | H | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | H | ОН |
| H | Н | Н | F | NH ₂ |
| H | Н | Н | F | NH-cyclopropyl |

| \mathbb{R}^1 | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | Y |
|----------------|----------------|----------------|-----------------|-----------------|
| H | H | Н | F | OH |
| H | Н | H | Cl | NH ₂ |
| H | H | H | Cl | NH-cyclopropyl |
| H | H | Н | Cl | OH |
| Н | Н | H | Br | NH ₂ |
| H | Н | Н | Br | NH-cyclopropyl |
| Н | Н | H | Br | OH |
| Н | Н | H | NH ₂ | NH ₂ |
| H | Н | Н | NH ₂ | NH-cyclopropyl |
| Н | H | H | NH ₂ | OH |
| H | H | Н | SH | NH ₂ |
| H | H | Н | SH | NH-cyclopropyl |
| H | H | H | SH | ОН |
| acetyl | H | H | Н | NH ₂ |
| acetyl | H | H | Н | NH-cyclopropyl |
| acetyl | Н | H | H | ОН |
| acetyl | H | H | F | NH ₂ |
| acetyl | H | H | F | NH-cyclopropyl |
| acetyl | Н | Н | F | ОН |
| Н | acetyl | acetyl | H | NH ₂ |
| H | acetyl | acetyl | H | NH-cyclopropyl |
| H | acetyl | acetyl | H | ОН |
| acetyl | acetyl | acetyl | H | NH ₂ |
| acetyl | acetyl | acetyl | Н | NH-cyclopropyl |
| acetyl | acetyl | acetyl | H | ОН |
| monophosphate | acetyl | acetyl | H | NH ₂ |
| monophosphate | acetyl | acetyl | H | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | H | ОН |
| diphosphate | acetyl | acetyl | H | NH ₂ |
| diphosphate | acetyl | acetyl | н | NH-cyclopropy |
| diphosphate | acetyl | acetyl | Н | OH |

| WO 01/90121 | | | | PCT/US01/1667 |
|----------------|----------------|----------------|----------------|-----------------|
| R ¹ | R ² | \mathbb{R}^3 | X ¹ | Y |
| triphosphate | acetyl | acetyl | H | NH ₂ |
| triphosphate | acetyl | acetyl | Н | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | Н | ОН |

Alternatively, the following nucleosides of Formula VII are prepared, using the appropriate sugar and pyrimidine or purine bases.

$$R^{1}O$$
 X
 R^{6}
 OR^{2}
 OR^{3}
 (VII)

wherein:

5

| R ¹ | \mathbb{R}^2 | R ³ | R ⁶ | X | Base |
|----------------|----------------|----------------|-----------------|---|-------------------|
| H | H | H | CH ₃ | 0 | 2,4-0- |
| | | | | | Diacetyluracil |
| H | Н | H | CH ₃ | 0 | Hypoxanthine |
| H | H | H | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthymine |
| H | H | H | CH ₃ | 0 | Thymine |
| H | H | H | CH ₃ | 0 | Cytosine |
| H | Н | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| H | H | H | CH ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| H | Н | H | CH ₃ | 0 | Uracil |
| H | H | H | CH ₃ | 0 | 5-Fluorouracil |
| H | H | Н | CH ₃ | S | 2,4-0- |
| | | | | | Diacetyluraci |
| H | H | Н | CH ₃ | S | Hypoxanthine |

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| \mathbb{R}^{1} | R ² | R ³ | R ⁶ | X | Base |
|------------------|----------------|----------------|-----------------|----|-------------------|
| H | Н | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetylthymine |
| H | Н | Н | CH ₃ | S | Thymine |
| Н | Н | H | CH ₃ | S | Cytosine |
| H | Н | Н | CH ₃ | S | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| Н | Н | Н | CH ₃ | S | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| H | Н | Н | CH ₃ | S. | Uracil |
| H | Н | Н | CH ₃ | S | 5-Fluorouracil |
| monophosphate | Н | Н | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetyluracil |
| monophosphate | H | Н | CH ₃ | 0 | Hypoxanthine |
| monophosphate | Н | H | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthym |
| monophosphate | Н | Н | CH ₃ | О | Thymine |
| monophosphate | Н | H . | CH ₃ | 0 | Cytosine |
| monophosphate | H | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| monophosphate | Н | H | CH ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| monophosphate | Н | Н | CH ₃ | 0 | Uracil |
| monophosphate | H | H | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | Н | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetyluracil |
| monophosphate | Н | Н | CH ₃ | S | Hypoxanthine |
| monophosphate | H | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetylthym |
| monophosphate | Н | Н | CH ₃ | S | Thymine |
| monophosphate | H | Н | CH ₃ | S | Cytosine |
| monophosphate | Н | H | CH ₃ | S | 4-(N-mono- |
| | | | | | acetyl)cytosine |

WO 01/90121 PCT/US01/16671 R⁶ $\overline{\mathbf{R}^1}$ $\overline{\mathbf{R^2}}$ \mathbb{R}^3 X Base monophosphate H H CH₃ S 4-(N,Ndiacetyl)cytosine CH₃ S monophosphate H H Uracil CH₃ S 5-Fluorouracil H H monophosphate H H CH₃ 2,4-0diphosphate O Diacetyluracil Hypoxanthine diphosphate H H CH₃ O CH₃ H O 2,4-0diphosphate H Diacetylthymine H CH₃ ō Thymine $\overline{\mathbf{H}}$ diphosphate CH₃ diphosphate H H O Cytosine 4-(N-mono- $\overline{\mathbf{H}}$ Η CH₃ 0 diphosphate acetyl)cytosine CH₃ O 4-(N,Ndiphosphate H H diacetyl)cytosine CH₃ diphosphate H H 0 Uracil H CH₃ 5-Fluorouracil diphosphate H 0 H S 2,4-0-H CH_3 diphosphate Diacetyluracil CH₃ S diphosphate Н H Hypoxanthine H CH₃ 2,4-0-H S diphosphate Diacetylthym CH₃ S diphosphate H H Thymine CH₃ diphosphate H H S Cytosine Н CH₃ O 2,4-0triphosphate H Diacetyluracil CH₃ Hypoxanthine $\overline{\mathbf{H}}$ H 0 triphosphate 2,4-0- \overline{H} H CH₃ $\overline{\mathbf{o}}$ triphosphate Diacetylthymine CH₃ H H Ō Thymine triphosphate H CH₃ O Cytosine triphosphate Η

| WO 01/90121 R ¹ | \mathbb{R}^2 | R ³ | R ⁶ | X | PCT/US01/16671 Base |
|-------------------------------|----------------|----------------|-----------------|---|----------------------|
| triphosphate | H | Н | CH ₃ | 0 | 4-(N-mono- |
| - | | | | | acetyl)cytosine |
| triphosphate | Н | H | CH ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| triphosphate | Н | Н | CH ₃ | 0 | Uracil |
| triphosphate | H | H | CH ₃ | 0 | 5-Fluorouracil |
| triphosphate | H | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetyluracil |
| triphosphate | H | Н | CH ₃ | S | Hypoxanthine |
| triphosphate | Н | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetylthymine |
| triphosphate | Н | Н | CH ₃ | S | Thymine |
| triphosphate | Н | Н | CH ₃ | S | Cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,4-0- |
| | | | | | Diacetyluracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Hypoxanthine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,4-0- |
| | | | | | Diacetylthymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Thymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Uracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 5-Fluorouracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,4-O- |
| | | | | | Diacetyluracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | Hypoxanthine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,4-O- |
| | | | | | Diacetylthymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | Thymine |

WO 01/90121 PCT/US01/16671 $\overline{\mathbf{R^6}}$ $\overline{\mathbf{R}^1}$ $\overline{\mathbf{R}^2}$ $\overline{\mathbf{R}^3}$ X Base monophosphate monophosphate monophosphate CF₃ S Cytosine 4-(N-mono-CF₃ S monophosphate monophosphate monophosphate acetyl)cytosine 4-(N,N-S monophosphate CF₃ monophosphate monophosphate diacetyl)cytosine Uracil CF₃ S monophosphate monophosphate monophosphate 5-Fluorouracil monophosphate monophosphate CF₃ S monophosphate CF₃ 4-(N,Nacetyl acetyl 0 acetyl diacetyl)cytosine 4-(N,N-S CF₃ acetyl acetyl acetyl diacetyl)cytosine 4-(N,N-2-bromo-0 acetyl acetyl acetyl diacetyl)cytosine vinyl 4-(N,N-2-bromoacetyl acetyl acetyi diacetyl)cytosine vinyl CH₃ 2-(N,N-diacetyl)-H Н H guanine 6-O-acetyl H H Н CH₃ O guanine H H CH₃ O 8-fluoroguanine Η CH_3 0 guanine H H Η H CH₃ $\overline{0}$ 6-(N,N-diacetyl)-H H adenine H CH₃ 2-fluoroadenine H 0 H CH₃ 8-fluoroadenine \overline{H} Η H 0 H H H CH₃ 0 2,8-difluoroadenine H CH₃ O adenine H $\overline{\mathbf{H}}$ H CH_3 S 2-(N,N-diacetyl)-H Η guanine \overline{s} 6-O-acetyl H CH₃ H H guanine

WO 01/90121 PCT/US01/16671 R⁶ $\overline{\mathbf{R}^1}$ $\overline{\mathbf{R^2}}$ R^3 X Base \overline{H} H S 8-fluoroguanine Η CH₃ H H Η CH₃ S guanine H S 6-(N,N-diacetyl)-H Η CH₃ adenine S 2-fluoroadenine Η H H CH₃ H CH₃ S 8-fluoroadenine H H CH₃ 2,8-difluoro-H H Η S adenine CH₃ S adenine H Н Н H CH₃ 0 2-(N,N-diacetyl)monophosphate Η guanine 6-O-acetyl H H CH₃ O monophosphate guanine CH₃ O 8-fluoroguanine H Η monophosphate Н CH₃ Ō Н guanine monophosphate 6-(N,N-diacetyl)-Η CH₃ O Η monophosphate adenine 2-fluoroadenine CH₃ $\overline{\mathbf{o}}$ Η Η monophosphate H Η CH_3 O 8-fluoroadenine monophosphate CH₃ 2,8-difluoro-H H O monophosphate adenine CH₃ H \overline{H} O adenine monophosphate H H CH₃ S 2-(N,N-diacetyl)monophosphate guanine 6-O-acetyl S H Η CH_3 monophosphate guanine 8-fluoroguanine H CH₃ S monophosphate H CH₃ S $\overline{\mathbf{H}}$ guanine Η monophosphate 6-(N,N-diacetyl)- CH_3 S monophosphate H Η adenine CH₃ H Η S 2-fluoroadenine monophosphate CH₃ Η S 8-fluoroadenine H monophosphate

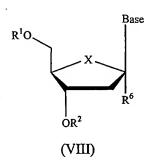
WO 01/90121 PCT/US01/16671 \mathbb{R}^1 R² \mathbb{R}^3 R⁶ X Base H CH₃ S 2,8-difluoromonophosphate H

| l II | ** | CII3 | | 2,0-41114010 |
|------|---------------------------------------|---------------------------------------|---|---|
| | | | | adenine |
| Н | Н | CH ₃ | S | adenine |
| H | H | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | guanine |
| Н | Н | CH ₃ | 0 | 6-O-acetyl |
| | | | | guanine |
| Н | Н | CH ₃ | 0 | 8-fluoroguanine |
| Н | Н | CH ₃ | 0 | guanine |
| Н | Н | CH ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | adenine |
| Н | Н | CH ₃ | 0 | 2-fluoroadenine |
| Н | Н | CH ₃ | 0 | 8-fluoroadenine |
| Н | Н | CH ₃ | 0 | 2,8-difluoro- |
| ĺ | | | | adenine |
| H | H | CH ₃ | 0 | adenine |
| Н | Н | CH ₃ | S | 2-(N,N-diacetyl)- |
| | | | | guanine |
| Н | Н | CH ₃ | S | 6-O-acetyl |
| | | | | guanine |
| Н | H | CH ₃ | S | 8-fluoroguanine |
| Н | Н | CH ₃ | S | guanine |
| H | H | CH ₃ | S | 6-(N,N-diacetyl)- |
| | | 7 | | adenine |
| H | H | CH ₃ | S | 2-fluoroadenine |
| Н | Н | CH ₃ | S | 8-fluoroadenine |
| Н | Н | CH ₃ | S | 2,8-difluoro- |
| | | | | adenine |
| H | Н | CH ₃ | S | adenine |
| Н | Н | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | guanine |
| | H H H H H H H H H H H H H H H H H H H | H H H H H H H H H H H H H H H H H H H | H H H CH3 H CH3 H H CH3 | H H H CH ₃ S H H H CH ₃ O H CH ₃ S H H H CH ₃ S S |

| triphosphate H H H CH3 CH3 O 6-O-acetyl guanine triphosphate H H H CH3 O 8-fluoroguanine triphosphate H H H CH3 O 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 | WO 01/90121 | | | | | PCT/US01/16671 |
|--|----------------|----------------|----------------|-----------------|---|-------------------|
| triphosphate H H H CH3 O S-fluoroguanine triphosphate H H H CH3 O G-(N,N-diacetyl)- adenine triphosphate H H H CH3 O S-fluoroadenine triphosphate H H CH3 CH3 O S-fluoroadenine triphosphate H H CH3 CH3 O S-fluoroadenine triphosphate H H CH3 CH3 S S-fluoroguanine triphosphate H H CH3 CH3 S S-fluoroguanine triphosphate H H CH3 CH3 S S-fluoroguanine triphosphate H H CH3 CH3 S S-fluoroadenine triphosphate M CH3 | R ¹ | R ² | \mathbb{R}^3 | R ⁶ | X | Base |
| triphosphate H H H CH3 O S-fluoroguanine triphosphate H H H CH3 O Guanine triphosphate H H H CH3 S Guanine triphosphate H H CH3 S Guanine Guan | triphosphate | Н | Н | CH ₃ | 0 | 6-O-acetyl |
| triphosphate H H H CH3 O guanine triphosphate H H H CH3 O G-(N,N-diacetyl)- adenine triphosphate H H H CH3 O 2-fluoroadenine triphosphate H H H CH3 O 3-fluoroadenine triphosphate H H H CH3 O 3-fluoroadenine triphosphate H H H CH3 O 2-ga-difluoro- adenine triphosphate H H H CH3 O 2-(N,N-diacetyl)- guanine triphosphate H H H CH3 S 6-O-acetyl guanine triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-guanine triphosphate H H CH3 S 3-guanine triphosphate H H CH3 S 3-guanine triphosphate H M CH3 S 3-fluoroadenine triphosphate H M CH3 S 3-guanine triphosphate H M CH3 S 3-guanine monophosphate monophosphate monophosphate CF3 O 3-guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine | | | | | | guanine |
| triphosphate H H H CH3 O 6-(N,N-diacetyl)- adenine triphosphate H H CH3 O 2-fluoroadenine triphosphate H H CH3 O 3-fluoroadenine triphosphate H H CH3 O 3-fluoroadenine triphosphate H H CH3 O 2-(N,N-diacetyl)- adenine triphosphate H H CH3 O 2-(N,N-diacetyl)- guanine triphosphate H H CH3 CH3 O 2-(N,N-diacetyl)- guanine triphosphate H H CH3 S 6-O-acetyl guanine triphosphate H H CH3 S 8-fluoroguanine triphosphate H H CH3 S S 8-fluoroguanine triphosphate H H CH3 S S 8-fluoroguanine triphosphate H H CH3 S S 6-(N,N-diacetyl)- adenine triphosphate H H CH3 S S 8-fluoroguanine triphosphate H H CH3 S S 8-fluoroadenine triphosphate H H CH3 S S 8-fluoroadenine triphosphate triphosphate H H CH3 S S S-fluoroadenine triphosphate triphosphate triphosphate H H CH3 S S S-fluoroadenine triphosphate triphosphate triphosphate H H CH3 S S S-fluoroadenine triphosphate triphosphate triphosphate H H CH3 S S S-fluoroadenine triphosphate triphosphate triphosphate H H CH3 S S S-fluoroadenine triphosphate triphosphate triphosphate triphosphate triphosphate H H CH3 S S S-fluoroadenine triphosphate tr | triphosphate | Н | Н | CH ₃ | 0 | 8-fluoroguanine |
| triphosphate H H H CH3 O 2-fluoroadenine triphosphate H H H CH3 O 8-fluoroadenine triphosphate H H H CH3 O 2,8-difluoro- adenine triphosphate H H H CH3 O 2,8-difluoro- adenine triphosphate H H H CH3 O 2-(N,N-diacetyl)- guanine triphosphate H H H CH3 S 8-fluoroguanine triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H CH3 CH3 S guanine triphosphate H H CH3 S guanine triphosphate H H CH3 S guanine triphosphate H H CH3 S 6-(N,N-diacetyl)- adenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 CH3 S 2,8-difluoro- adenine triphosphate M M CH3 S 3-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine | triphosphate | H | H | CH ₃ | 0 | guanine |
| triphosphate H H H CH3 CH3 O 2-fluoroadenine triphosphate H H H CH3 S 6-O-acetyl guanine triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 CH3 S 2-fluoroadenine triphosphate H H CH3 CH3 S 3-fluoroadenine triphosphate H H CH3 CH3 S 3-fluoroadenine triphosphate H CH3 CH3 S 3-fluoroadenine triphosphate H CH3 CH3 S 3-fluoroadenine triphosphate M CH3 S 3-fluoroadenine | triphosphate | Н | Н | CH ₃ | 0 | 6-(N,N-diacetyl)- |
| triphosphate H H H CH3 O 8-fluoroadenine triphosphate H H H CH3 O 2,8-difluoro- adenine triphosphate H H H CH3 O 2-(N,N-diacetyl)- guanine triphosphate H H H CH3 S 8-fluoroguanine triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H CH3 S 3-fluoroguanine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 CH3 S 8-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 CH3 S 3-fluoroadenine triphosphate Monophosphate Monophosphate CF3 O 3-coetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine | | | | | | adenine |
| triphosphate H H H CH3 S 3-denine triphosphate H H H CH3 S 3-denome triphosphate H H CH3 S 3-denome triphosphate H H CH3 CH3 S 3-denome triphosphate H H CH3 CH3 S 3-denome triphosphate H H CH3 S 3-denome triphosphate H H CH3 CH3 S 3-denome triphosphate H H CH3 CH3 S 3-denome triphosphate H H CH3 CH3 S 3-denome triphosphate M H CH3 CF3 O 3-denome monophosphate M Monophosphate CF3 O 6-O-acetyl guanine monophosphate M Monophosphate M Monophosphate CF3 O 8-fluoroguanine monophosphate M Monophosphate M Monophosphate CF3 O 8-fluoroguanine | triphosphate | Н | Н | CH ₃ | 0 | 2-fluoroadenine |
| triphosphate H H H CH3 S 2-fluoroadenine triphosphate H H H CH3 S 3-fluoroadenine triphosphate H H H CH3 S 2-fluoroadenine triphosphate H H H CH3 S 2-fluoroadenine triphosphate H H H CH3 S 3-fluoroadenine triphosphate H H H CH3 S 2-fluoroadenine triphosphate H H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine | triphosphate | H | H | CH ₃ | 0 | 8-fluoroadenine |
| triphosphate triphosphate triphosphate H H H CH3 CH3 CH3 CH3 CH3 CH3 | triphosphate | Н | Н | CH ₃ | 0 | 2,8-difluoro- |
| triphosphate H H H CH3 S guanine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 3-fluoroadenine triphosphate H H CH3 S 6-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | | | | | | adenine |
| triphosphate H H CH3 S 6-O-acetyl guanine triphosphate H H CH3 S 8-fluoroguanine triphosphate H H CH3 S guanine Triphosphate H H CH3 S G-(N,N-diacetyl)- adenine triphosphate H H CH3 S S-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine Triphosphate H H CH3 S 8-fluoroadenine Triphosphate H H CH3 S 8-fluoroadenine Triphosphate Triphosphate H H CH3 S 8-fluoroadenine Triphosphate Triphosphate Triphosphate Triphosphate H H CH3 S Adenine Triphosphate Tri | triphosphate | H | H | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| triphosphate H H H CH3 S guanine triphosphate H H H CH3 S guanine triphosphate H H CH3 CH3 S guanine triphosphate H H CH3 CH3 S G-(N,N-diacetyl)- adenine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate H M CH3 S adenine monophosphate monophosphate Monophosphate CF3 O C-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O S-fluoroguanine monophosphate monophosphate monophosphate CF3 O S-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | | | | | | guanine |
| triphosphate H H H CH3 S guanine triphosphate H H CH3 S guanine triphosphate H H CH3 S guanine triphosphate H H CH3 S G-(N,N-diacetyl)- adenine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate H monophosphate CH3 S adenine monophosphate monophosphate Monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate Monophospha | triphosphate | Н | H | CH ₃ | S | 6-O-acetyl |
| triphosphate H H H CH3 S guanine triphosphate H H H CH3 S 6-(N,N-diacetyl)- adenine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate H H CH3 S adenine triphosphate H H CH3 S adenine monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | | | | | | guanine |
| triphosphate H H CH3 S 6-(N,N-diacetyl)- adenine triphosphate H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | triphosphate | Н | Н | CH ₃ | S | 8-fluoroguanine |
| triphosphate H H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate H H CH3 S adenine triphosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | triphosphate | Н | Н | CH ₃ | S | guanine |
| triphosphate H H H CH3 S 2-fluoroadenine triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | triphosphate | Н | Н | CH ₃ | S | 6-(N,N-diacetyl)- |
| triphosphate H H CH3 S 8-fluoroadenine triphosphate H H CH3 CH3 S 2,8-difluoro- adenine triphosphate H H CH3 S adenine triphosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | | | _ | | | adenine |
| triphosphate H H H CH3 S adenine triphosphate H H CH3 S adenine monophosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | triphosphate | Н | Н | CH ₃ | S | 2-fluoroadenine |
| triphosphate H H CH3 S adenine monophosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | triphosphate | Н | Н | CH ₃ | S | 8-fluoroadenine |
| triphosphate H H CH3 S adenine monophosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate monophosphate CF3 O guanine | triphosphate | Н | H | CH ₃ | S | 2,8-difluoro- |
| monophosphate monophosphate monophosphate CF3 O 2-(N,N-diacetyl)- guanine monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate cF3 O guanine | | | | _ | | adenine |
| monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate CF3 O guanine | triphosphate | Н | Н | CH ₃ | S | adenine |
| monophosphate monophosphate monophosphate CF3 O 6-O-acetyl guanine monophosphate monophosphate monophosphate CF3 O 8-fluoroguanine monophosphate monophosphate CF3 O guanine | monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2-(N,N-diacetyl)- |
| monophosphate monophosphate monophosphate CF ₃ O guanine monophosphate monophosphate CF ₃ O guanine | | | | | | guanine |
| monophosphate monophosphate monophosphate CF ₃ O 8-fluoroguanine monophosphate monophosphate CF ₃ O guanine | monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 6-O-acetyl |
| monophosphate monophosphate CF ₃ O guanine | | | | | | guanine |
| | monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroguanine |
| monophosphate monophosphate CF ₃ O 6-(N,N-diacetyl)- | monophosphate | monophosphate | monophosphate | CF ₃ | 0 | guanine |
| | monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 6-(N,N-diacetyl)- |
| adenine | | | | | | |
| monophosphate monophosphate CF ₃ O 2-fluoroadenine | monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2-fluoroadenine |

| WO 01/90121 | \mathbb{R}^2 | \mathbb{R}^3 | R ⁶ | X | PCT/US01/16671 Base |
|---------------|----------------|----------------|-----------------|---|------------------------|
| | | | | | |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2-(N,N-diacetyl)- |
| | | | - | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 8-fluoroguanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 8-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | adenine |
| acetyl | acetyl | acetyl | CF ₃ | 0 | guanine |
| acetyl | acetyl | acetyl | CF ₃ | S | guanine |
| acetyl | acetyl | acetyl | 2-bromo- | 0 | guanine |
| | | | vinyl | | |
| acetyl | acetyl | acetyl | 2-bromo- | S | guanine |
| | | | vinyl | | |

Alternatively, the following nucleosides of Formula VIII are prepared, using the appropriate sugar and pyrimidine or purine bases.



| R ¹ | \mathbb{R}^2 | R ⁶ | X | Base |
|----------------|----------------|-----------------|---|---------------------------|
| Н | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| H | Н | CH ₃ | 0 | Hypoxanthine |
| Н | H | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| H | H | CH ₃ | 0 | Thymine |
| H | H | CH ₃ | 0 | Cytosine |
| H | Н | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| Н | Н | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| Н | Н | CH ₃ | 0 | Uracil |
| Н | Н | CH ₃ | 0 | 5-Fluorouracil |
| H | Н | CH ₃ | S | 2,4-O-Diacetyluracil |
| H | H | CH ₃ | S | Hypoxanthine |
| Н | H | CH ₃ | S | 2,4-O-Diacetylthymine |
| H | Н | CH ₃ | S | Thymine |
| Н | Н | CH ₃ | S | Cytosine |
| Н | H | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| Н | Н | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| H | H | CH ₃ | S | Uracil |
| Н | H | CH ₃ | S | 5-Fluorouracil |
| monophosphate | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | H | CH ₃ | 0 | Hypoxanthine |
| monophosphate | H | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| monophosphate | Н | CH ₃ | 0 | Thymine |
| monophosphate | Н | CH ₃ | 0 | Cytosine |
| monophosphate | H | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | H | CH ₃ | 0 | Uracil |
| monophosphate | Н | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | Н | CH ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | Н | CH ₃ | S | Hypoxanthine |
| monophosphate | H | CH ₃ | S | 2,4-O-Diacetylthymine |
| L | 1 | 1 | | |

| R ¹ | $\mathbf{\dot{R}^2}$ | ~ h | | |
|----------------|----------------------|-----------------|----|---------------------------|
| | IX | R ⁶ | X | Base |
| monophosphate | H | CH ₃ | S | Thymine |
| monophosphate | Н | CH ₃ | S | Cytosine |
| monophosphate | Н | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | Н | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | H | CH ₃ | S | Uracil |
| monophosphate | Н | CH ₃ | S | 5-Fluorouracil |
| diphosphate | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| diphosphate | H | CH ₃ | 0 | Hypoxanthine |
| diphosphate | H | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| diphosphate | H | CH ₃ | 0 | Thymine |
| diphosphate | Н | CH ₃ | 0 | Cytosine |
| diphosphate | H | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| diphosphate | Н | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| diphosphate | Н | CH ₃ | 0 | Uracil |
| diphosphate | Н | CH ₃ | 0 | 5-Fluorouracil |
| diphosphate | Н | CH ₃ | S | 2,4-O-Diacetyluracil |
| diphosphate | Н | CH ₃ | S | Hypoxanthine |
| diphosphate | Н | CH ₃ | S | 2,4-O-Diacetylthymine |
| diphosphate | Н | CH ₃ | S | Thymine |
| diphosphate | Н | CH ₃ | S | Cytosine |
| diphosphate | Н | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| diphosphate | Н | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| diphosphate | Н | CH ₃ | S | Uracil |
| diphosphate | Н | CH ₃ | S | 5-Fluorouracil |
| triphosphate | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| triphosphate | Н | CH ₃ | 0 | Hypoxanthine |
| triphosphate | H | CH ₃ | 0 | 2,4-O-diacethylthymine |
| triphosphate | H | CH ₃ | 0 | Thymine |
| triphosphate | H | CH ₃ | Ο. | Cytosine |
| triphosphate | Н | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| triphosphate | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |

| WO 01/90121 | | | | PCT/US01/16671 |
|---------------|----------------|-----------------|---|---------------------------|
| R¹ | \mathbb{R}^2 | \mathbb{R}^6 | X | Base |
| triphosphate | H | CH ₃ | 0 | Uracil |
| triphosphate | Н | CH₃ | 0 | 5-Fluorouracil |
| triphosphate | H | CH ₃ | S | 2,4-O-Diacetyluracil |
| triphosphate | H | CH ₃ | S | Hypoxanthine |
| triphosphate | H | CH ₃ | S | 2,4-O-Diacetylthymine |
| triphosphate | Н | CH ₃ | S | Thymine |
| triphosphate | H | CH ₃ | S | Cytosine |
| triphosphate | H | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| triphosphate | H | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| triphosphate | Н | CH ₃ | S | Uracil |
| triphosphate | Н | CH ₃ | S | 5-Fluorouracil |
| monophosphate | monophosphate | CF ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | monophosphate | CF ₃ | 0 | Hypoxanthine |
| monophosphate | monophosphate | CF ₃ | 0 | 2,4-O-Diacetylthymine |
| monophosphate | monophosphate | CF ₃ | 0 | Thymine |
| monophosphate | monophosphate | CF ₃ | 0 | Cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | Uracil |
| monophosphate | monophosphate | CF ₃ | 0 | 5-Fluorouracil |
| monophosphate | monophosphate | CF ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | monophosphate | CF ₃ | S | Hypoxanthine |
| monophosphate | monophosphate | CF ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | monophosphate | CF ₃ | S | Thymine |
| monophosphate | monophosphate | CF ₃ | S | Cytosine |
| monophosphate | monophosphate | CF ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | S | Uracil |
| monophosphate | monophosphate | CF ₃ | S | 5-Fluorouracil |
| acetyl | acetyl | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| acetyl | acetyl | CF ₃ | S | 4-(N,N-diacetyl)cytosine |

| R ¹ | \mathbb{R}^2 | R ⁶ | X | PCT/US01/16671 Base |
|----------------|----------------|-----------------|---|-----------------------------|
| acetyl | acetyl | 2-bromo- | 0 | 4-(N,N-diacetyl)cytosine |
| acciyi | acctyr | vinyl | | 4-(14,14-411acoty1)cytosino |
| | | | | 4-(N,N-diacetyl)cytosine |
| acetyl | acetyl | 2-bromo- | S | 4-(11,11-diacetyl)cytosine |
| | | vinyl | J | |
| H | H | CH ₃ | 0 | 2-(N,N-diacetyl)-guanine |
| H | H | CH ₃ | 0 | 6-O-acetyl guanine |
| H | H | CH ₃ | 0 | 8-fluoroguanine |
| H | Н | CH ₃ | 0 | guanine |
| H . | Н | CH ₃ | О | 6-(N,N-diacetyl)-adenine |
| Н | H | CH ₃ | 0 | 2-fluoroadenine |
| H | H | CH ₃ | 0 | 8-fluoroadenine |
| H | Н | CH ₃ | 0 | 2,8-difluoro-adenine |
| H | H | CH ₃ | 0 | adenine |
| Н | H | CH ₃ | S | 2-(N,N-diacetyl)-guanine |
| Н | Н | CH ₃ | S | 6-O-acetyl guanine |
| H | Н | CH ₃ | S | 8-fluoroguanine |
| H | Н | CH ₃ | S | guanine |
| H | Н | CH ₃ | S | 6-(N,N-diacetyl)-adenine |
| H | H | CH ₃ | S | 2-fluoroadenine |
| H | Н | CH ₃ | S | 8-fluoroadenine |
| Н | H | CH ₃ | S | 2,8-difluoro-adenine |
| Н | Н | CH ₃ | S | adenine |
| monophosphate | H | CH ₃ | 0 | 2-(N,N-diacetyl)-guanine |
| monophosphate | H | CH ₃ | 0 | 6-O-acetyl guanine |
| monophosphate | Н | CH ₃ | 0 | 8-fluoroguanine |
| monophosphate | Н | CH ₃ | 0 | guanine |
| monophosphate | H | CH ₃ | 0 | 6-(N,N-diacetyl)-adenine |
| monophosphate | Н | CH ₃ | 0 | 2-fluoroadenine |
| monophosphate | Н | CH ₃ | 0 | 8-fluoroadenine |
| monophosphate | H | CH ₃ | 0 | 2,8-difluoro-adenine |
| monophosphate | H | CH ₃ | 0 | adenine |

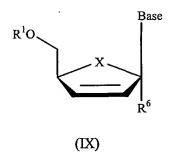
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|-------------|----------------|
| | |

| R ¹ | \mathbb{R}^2 | R ⁶ | X | PCT/US01/16671 Base |
|----------------|----------------|-----------------|---|--------------------------|
| monophosphate | Н | CH ₃ | S | 2-(N,N-diacetyl)-guanine |
| monophosphate | Н | CH ₃ | S | 6-O-acetyl guanine |
| monophosphate | H | CH ₃ | S | 8-fluoroguanine |
| monophosphate | Н | CH ₃ | S | guanine |
| monophosphate | Н | CH ₃ | S | 6-(N,N-diacetyl)-adenine |
| monophosphate | Н | CH ₃ | S | 2-fluoroadenine |
| monophosphate | Н | CH ₃ | S | 8-fluoroadenine |
| monophosphate | Н | CH ₃ | S | 2,8-difluoro-adenine |
| monophosphate | Н | CH ₃ | S | adenine |
| diphosphate | Н | CH ₃ | 0 | 2-(N,N-diacetyl)-guanine |
| diphosphate | Н | CH ₃ | 0 | 6-O-acetyl guanine |
| diphosphate | H | CH ₃ | 0 | 8-fluoroguanine |
| diphosphate | Н | CH ₃ | 0 | guanine |
| diphosphate | Н | CH ₃ | 0 | 6-(N,N-diacetyl)-adenine |
| diphosphate | H | CH ₃ | 0 | 2-fluoroadenine |
| diphosphate | Н | CH ₃ | 0 | 8-fluoroadenine |
| diphosphate | H | CH ₃ | 0 | 2,8-difluoro-adenine |
| diphosphate | Н | CH ₃ | 0 | adenine |
| diphosphate | H | CH ₃ | S | 2-(N,N-diacetyl)-guanine |
| diphosphate | Н | CH ₃ | S | 6-O-acetyl guanine |
| diphosphate | H | CH ₃ | S | 8-fluoroguanine |
| diphosphate | Н | CH ₃ | S | guanine |
| diphosphate | H | CH ₃ | S | 6-(N,N-diacetyl)-adenine |
| diphosphate | H | CH ₃ | S | 2-fluoroadenine |
| diphosphate | H | CH ₃ | S | 8-fluoroadenine |
| diphosphate | Н | CH ₃ | S | 2,8-difluoro-adenine |
| diphosphate | H | CH ₃ | S | adenine |
| triphosphate | H | CH ₃ | 0 | 2-(N,N-diacetyl)-guanine |
| triphosphate | H | CH ₃ | 0 | 6-O-acetyl guanine |
| triphosphate | H | CH ₃ | 0 | 8-fluoroguanine |
| triphosphate | Н | CH ₃ | 0 | guanine |

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|-------------------------------|----------------|-----------------|-----|--------------------------|
| WO 01/90121 R ¹ | \mathbb{R}^2 | R ⁶ | X | Base |
| triphosphate | Н | CH ₃ | 0 | 6-(N,N-diacetyl)-adenine |
| triphosphate | Н | CH ₃ | 0 | 2-fluoroadenine |
| triphosphate | Н | CH ₃ | 0 | 8-fluoroadenine |
| triphosphate | Н | CH ₃ | 0 | 2,8-difluoro-adenine |
| triphosphate | Н | CH ₃ | 0 | adenine |
| triphosphate | H | CH ₃ | S | 2-(N,N-diacetyl)-guanine |
| triphosphate | H | CH ₃ | S | 6-O-acetyl guanine |
| triphosphate | Н | CH ₃ | S | 8-fluoroguanine |
| triphosphate | H | CH ₃ | S | guanine |
| triphosphate | Н . | CH ₃ | S | 6-(N,N-diacetyl)-adenine |
| triphosphate | Н | CH ₃ | S | 2-fluoroadenine |
| triphosphate | Н | CH ₃ | S | 8-fluoroadenine |
| triphosphate | H | CH ₃ | S · | 2,8-difluoro-adenine |
| triphosphate | H | CH ₃ | S | adenine |
| monophosphate | monophosphate | CF ₃ | 0 | 2-(N,N-diacetyl)-guanine |
| monophosphate | monophosphate | CF ₃ | 0 | 6-O-acetyl guanine |
| monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroguanine |
| monophosphate | monophosphate | CF ₃ | 0 | guanine |
| monophosphate | monophosphate | CF ₃ | 0 | 6-(N,N-diacetyl)-adenine |
| monophosphate | monophosphate | CF ₃ | 0 | 2-fluoroadenine |
| monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroadenine |
| monophosphate | monophosphate | CF ₃ | 0 | 2,8-difluoro-adenine |
| monophosphate | monophosphate | CF ₃ | 0 | adenine |
| monophosphate | monophosphate | CF ₃ | S | 2-(N,N-diacetyl)-guanine |
| monophosphate | monophosphate | CF ₃ | S | 6-O-acetyl guanine |
| monophosphate | monophosphate | CF ₃ | S | 8-fluoroguanine |
| monophosphate | monophosphate | CF ₃ | S | guanine |
| monophosphate | monophosphate | CF ₃ | S | 6-(N,N-diacetyl)-adenine |
| monophosphate | monophosphate | CF ₃ | S | 2-fluoroadenine |
| monophosphate | monophosphate | CF ₃ | S | 8-fluoroadenine |
| monophosphate | monophosphate | CF ₃ | S | 2,8-difluoro-adenine |

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|----------------|----------------|-------------------|---|----------------|
| \mathbb{R}^1 | R ² | R ⁶ | X | Base |
| monophosphate | monophosphate | CF ₃ | S | adenine |
| acetyl | acetyl | CF ₃ | 0 | guanine |
| acetyl | acetyl | CF ₃ | S | guanine |
| acetyl | acetyl | 2-bromo- vinyl | 0 | guanine |
| acetyl | acetyl | 2-bromo- vinyl | S | guanine |

Alternatively, the following nucleosides of Formula IX are prepared, using the appropriate sugar and pyrimidine or purine bases.



wherein:

| R ¹ | R ⁶ | X | Base | | |
|----------------|-----------------|---|---------------------------|--|--|
| Н | CH ₃ | 0 | 2,4-O-Diacetyluracil | | |
| H | CH ₃ | 0 | Hypoxanthine | | |
| H | CH ₃ | 0 | 2,4-O-Diacetylthymine | | |
| Н | CH ₃ | 0 | Thymine | | |
| H | CH ₃ | 0 | Cytosine | | |
| Н | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine | | |
| Н | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine | | |
| H | CH ₃ | 0 | Uracil | | |
| H | CH ₃ | 0 | 5-Fluorouracil | | |
| H | CH ₃ | S | 2,4-O-Diacetyluracil | | |
| H | CH ₃ | S | Hypoxanthine | | |
| H | CH ₃ | S | 2,4-O-Diacetylthymine | | |

| WO 01/90121 R ¹ | R ⁶ | Tv | Pers |
|-------------------------------|-----------------|-----|---------------------------|
| | | X | Base |
| Н | CH₃ | S | Thymine |
| Н | CH ₃ | S | Cytosine |
| Н | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| H | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| Н | CH ₃ | S | Uracil |
| H | CH ₃ | S | 5-Fluorouracil |
| monophosphate | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | CH ₃ | 0 | Hypoxanthine |
| monophosphate | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| monophosphate | CH ₃ | 0 | Thymine |
| monophosphate | CH ₃ | 0 | Cytosine |
| monophosphate | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | CH ₃ | 0 | Uracil |
| monophosphate | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | CH ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | CH ₃ | S | Hypoxanthine |
| monophosphate | CH ₃ | S · | 2,4-O-Diacetylthymine |
| monophosphate | CH ₃ | S | Thymine |
| monophosphate | CH ₃ | S | Cytosine |
| monophosphate | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | CH ₃ | S | 4-(N,N-diacetyl)cytos |
| monophosphate | CH ₃ | S | Uracil |
| monophosphate | CH ₃ | S | 5-Fluorouracil |
| diphosphate | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| diphosphate | CH ₃ | 0 | Hypoxanthine |
| diphosphate | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| diphosphate | CH ₃ | 0 | Thymine |
| diphosphate | CH ₃ | 0 | Cytosine |
| diphosphate | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| diphosphate | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| L | _L | | |

| WO 01/90121 R ¹ R ⁶ X Base | | | | | |
|--|---|---|--|--|--|
| R ⁶ | X | Base | | | |
| CH ₃ | 0 | Uracil | | | |
| CH ₃ | 0 | 5-Fluorouracil | | | |
| CH ₃ | S | 2,4-O-Diacetyluracil | | | |
| CH ₃ | S | Hypoxanthine | | | |
| CH ₃ | S | 2,4-O-Diacetylthymine | | | |
| CH ₃ | S | Thymine | | | |
| CH ₃ | S | Cytosine | | | |
| CH ₃ | 0 | 2,4-O-Diacetyluracil | | | |
| CH ₃ | 0 | Hypoxanthine | | | |
| CH ₃ | 0 | 2,4-O-Diacetylthymine | | | |
| CH ₃ | 0 | Thymine | | | |
| CH ₃ | 0 | Cytosine | | | |
| CH ₃ | 0 | 4-(N-mono-acetyl)cytosine | | | |
| CH ₃ | 0 | 4-(N,N-diacetyl)cytosine | | | |
| CH ₃ | 0 | Uracil | | | |
| CH ₃ | 0 | 5-Fluorouracil | | | |
| CH ₃ | S | 2,4-O-Diacetyluracil | | | |
| CH ₃ | S | Hypoxanthine | | | |
| CH ₃ | S | 2,4-O-Diacetylthymine | | | |
| CH ₃ | S | Thymine | | | |
| CH ₃ | S | Cytosine | | | |
| CF ₃ | 0 | 2,4-O-Diacetyluracil | | | |
| CF ₃ | 0 | Hypoxanthine | | | |
| CF ₃ | 0 | 2,4-O-Diacetylthymine | | | |
| CF ₃ | . 0 | Thymine | | | |
| CF ₃ | 0 | Cytosine | | | |
| CF ₃ | 0 | 4-(N-mono-acetyl)cytosine | | | |
| CF ₃ | 0 | 4-(N,N-diacetyl)cytos | | | |
| CF ₃ | 0 | Uracil | | | |
| CF ₃ | 0 | 5-Fluorouracil | | | |
| CF ₃ | S | 2,4-O-Diacetyluracil | | | |
| | CH ₃ CG ₃ | CH3 O CH3 O CH3 S CH3 S CH3 S CH3 S CH3 O CH3 S CF3 O CF3 O CF3 O CF3 O CF3 O CF3 O CF3 O | | | |

| R ¹ | R ⁶ | X | Base |
|----------------|-----------------|---|---------------------------|
| monophosphate | CF ₃ | S | Hypoxanthine |
| monophosphate | CF ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | CF ₃ | S | Thymine |
| monophosphate | CF ₃ | S | Cytosine |
| monophosphate | CF ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | CF ₃ | S | Uracil |
| monophosphate | CF ₃ | S | 5-Fluorouracil |
| acetyl | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| acetyl | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| acetyl | 2-bromo-vinyl | 0 | 4-(N,N-diacetyl)cytosine |
| acetyl | 2-bromo-vinyl | S | 4-(N,N-diacetyl)cytosine |

Alternatively, the following nucleosides of Formula XVI are prepared, using the appropriate sugar and pyrimidine or purine bases.

$$R^{10}$$
 X R^{8} R^{6} R^{9} R^{7} R^{7}

wherein:

| R ¹ | R ⁶ | \mathbb{R}^7 | R ⁸ | X | Base | R ¹⁰ | R ⁹ |
|----------------|-----------------|----------------|----------------|---|---------------------------|-----------------|----------------|
| H | CH ₃ | Н | Н | 0 | 2,4-O-Diacetyluracil | ОН | Me |
| Н | CH ₃ | Н | H | 0 |) Hypoxanthine | | Me |
| Н | CH ₃ | Н | Н | 0 | 2,4-O-Diacetylthymine | | Me |
| H | CH ₃ | H | H | 0 | Thymine | ОН | Me |
| Н | CH ₃ | Н | H | 0 | Cytosine | ОН | Me |
| H | CH ₃ | H | Н | 0 | 4-(N-mono-acetyl)cytosine | ОН | Me |
| Н | CH ₃ | H | H | 0 | 4-(N,N-diacetyl)cytosine | ОН | Me |

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|----------------------------|-----------------|----------------|----------------|---|---------------------------|-----------------|----------------|
| R ¹ | R ⁶ | \mathbb{R}^7 | R ⁸ | X | Base | R ¹⁰ | R ⁹ |
| H | CH ₃ | Н | H | 0 | Uracil | OH | Me |
| H | CH ₃ | H | H | 0 | 5-Fluorouracil | OH | Me |
| Н | CH ₃ | H | Н | S | 2,4-O-Diacetyluracil | ОН | Me |
| Н | CH ₃ | Н | H | S | Hypoxanthine | OH | Me |
| H | CH ₃ | Н | Н | S | 2,4-O-Diacetylthymine | ОН | Me |
| Н | CH ₃ | H | Н | S | Thymine | ОН | Me |
| Н | CH ₃ | H | H | S | Cytosine | OH | Me |
| Н | CH ₃ | H | H | S | 4-(N-mono-acetyl)cytosine | ОН | Me |
| Н | CH ₃ | Н | Н | S | 4-(N,N-diacetyl)cytosine | OH | Me |
| H | CH ₃ | Н | Н | S | Uracil | OH | Me |
| Н | CH ₃ | H | H | S | 5-Fluorouracil | OH | Me |
| monophosphate | CH ₃ | Н | Н | 0 | 2,4-O-Diacetyluracil | OH | Me |
| monophosphate | CH ₃ | H | Н | 0 | Hypoxanthine | ОН | Me |
| monophosphate | CH ₃ | H | H | Ö | 2,4-O-Diacetylthymine | OH | Me |
| monophosphate | CH ₃ | Н | Н | 0 | Thymine | OH | Me |
| monophosphate | CH ₃ | H | Н | 0 | Cytosine | OH | Me |
| monophosphate | CH ₃ | Н | Н | 0 | 4-(N-mono-acetyl)cytosine | ОН | Me |
| monophosphate | CH ₃ | Н | H | 0 | 4-(N,N-diacetyl)cytosine | ОН | Me |
| monophosphate | CH ₃ | Н | Н | 0 | Uracil | OH | Me |
| monophosphate | CH ₃ | Н | Н | 0 | 5-Fluorouracil | OH | Me |
| monophosphate | CH ₃ | Н | H | S | 2,4-O-Diacetyluracil | OH | Me |
| monophosphate | CH ₃ | Н | Н | S | Hypoxanthine | OH | Me |
| monophosphate | CH ₃ | H | H | S | 2,4-O-Diacetylthymine | OH | Me |
| monophosphate | CH ₃ | Н | H | S | Thymine | OH | Me |
| monophosphate | CH ₃ | H | H | S | Cytosine | OH | Me |
| monophosphate | CH ₃ | Н | H | S | 4-(N-mono-acetyl)cytosine | OH | Me |
| monophosphate | CH ₃ | Н | H | S | 4-(N,N-diacetyl)cytosine | OH | Me |
| monophosphate | CH ₃ | Н | H | S | Uracil | ОН | Me |
| monophosphate | CH ₃ | Н | H | S | 5-Fluorouracil | OH | Me |
| diphosphate | CH ₃ | H | H | 0 | 2,4-O-Diacetyluracil | OH | Me |
| diphosphate | CH ₃ | Н | H | 0 | Hypoxanthine | ОН | Me |
| L | | | | | | | |

| R ¹ | \mathbb{R}^6 | \mathbf{R}^7 | R ⁸ | X | Base | R ¹⁰ | R ⁹ |
|----------------|-----------------|----------------|----------------|----|---------------------------|-----------------|----------------|
| diphosphate | CH ₃ | H | H | 0 | 2,4-O-Diacetylthymine | ОН | Me |
| diphosphate | CH ₃ | H | Н | 0 | Thymine | ОН | Me |
| diphosphate | CH ₃ | Н | Н | 0 | Cytosine | ОН | Me |
| diphosphate | CH ₃ | H | Н | 0 | 4-(N-mono-acetyl)cytosine | OH | Me |
| diphosphate | CH ₃ | H | Н | 0 | 4-(N,N-diacetyl)cytosine | OH | Me |
| diphosphate | CH ₃ | H | H | 0 | Uracil | ОН | Me |
| diphosphate | CH ₃ | Н | H | 0 | 5-Fluorouracil | ОН | Me |
| diphosphate | CH ₃ | Н | H | S | 2,4-O-Diacetyluracil | OH | Me |
| diphosphate | CH ₃ | Н | Н | S | Hypoxanthine | OH | Me |
| diphosphate | CH ₃ | Н | Н | S | 2,4-O-Diacetylthymine | ОН | Me |
| diphosphate | CH ₃ | Н | Н | ·S | Thymine | ОН | Me |
| diphosphate | CH ₃ | Н | H | S | Cytosine | OH | Me |
| triphosphate | CH ₃ | Н | Н | 0 | 2,4-O-Diacetyluracil | ОН | Me |
| triphosphate | CH ₃ | H | Н | 0 | Hypoxanthine | ОН | Me |
| triphosphate | CH ₃ | Н | Н | 0 | 2,4-O-Diacetylthymine | ОН | Me |
| triphosphate | CH ₃ | H | Н | 0 | Thymine | ОН | Me |
| triphosphate | CH ₃ | H | H | 0 | Cytosine | ОН | Me |
| triphosphate | CH ₃ | Н | Н | 0 | 4-(N-mono-acetyl)cytosine | ОН | Me |
| triphosphate | CH ₃ | Н | Н | 0 | 4-(N,N-diacetyl)cytosine | ОН | Me |
| triphosphate | CH ₃ | Н | Н | 0 | Uracil | OH | Me |
| triphosphate | CH ₃ | Н | Н | 0 | 5-Fluorouracil | OH | Me |
| triphosphate | CH ₃ | Н | Н | S | 2,4-O-Diacetyluracil | OH | Me |
| triphosphate | CH ₃ | Н | Н | S | Hypoxanthine | OH | Me |
| triphosphate | CH ₃ | H | Н | S | 2,4-O-Diacetylthymine | ОН | Me |
| triphosphate | CH ₃ | H | Н | S | Thymine | ОН | Me |
| triphosphate | CH ₃ | Н | Н | S | Cytosine | OH | Me |
| monophosphate | CF ₃ | Н | Н | 0 | 2,4-O-Diacetyluracil | OH | Me |
| monophosphate | CF ₃ | Н | Н | 0 | Hypoxanthine | ОН | Me |
| monophosphate | CF ₃ | Н | H | 0 | 2,4-O-Diacetylthymine OH | | Me |
| monophosphate | CF ₃ | Н | H | 0 | Thymine C | | Me |
| monophosphate | CF ₃ | H | H | 0 | Cytosine | ОН | Me |

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|----------------|-----------------|----------------|----------------|---|---------------------------|-----------------|----------------|--|
| \mathbb{R}^1 | R ⁶ | R ⁷ | R ⁸ | X | Base | R ¹⁰ | R ⁹ | |
| monophosphate | CF ₃ | Н | Н | 0 | 4-(N-mono-acetyl)cytosine | OH | Me | |
| monophosphate | CF ₃ | H | H | 0 | 4-(N,N-diacetyl)cytosine | OH | Me | |
| monophosphate | CF ₃ | Н | H | 0 | Uracil | OH | Me | |
| monophosphate | CF ₃ | Н | H | 0 | 5-Fluorouracil | OH | Me | |
| monophosphate | CF ₃ | Н | Н | S | 2,4-O-Diacetyluracil | ОН | Me | |
| monophosphate | CF ₃ | Н | H | S | Hypoxanthine | OH | Me | |
| monophosphate | CF ₃ | H | Н | S | 2,4-O-Diacetylthymine | ОН | Me | |
| monophosphate | CF ₃ | Н | Н | S | Thymine | OH | Me | |
| monophosphate | CF ₃ | H | H | S | Cytosine | ОН | Me | |
| monophosphate | CF ₃ | H | H | S | 4-(N-mono-acetyl)cytosine | ОН | Me | |
| monophosphate | CF ₃ | Н | H | S | 4-(N,N-diacetyl)cytosine | ОН | Me | |
| monophosphate | CF ₃ | Н | H | S | Uracil | ОН | Me | |
| monophosphate | CF ₃ | H | H | S | 5-Fluorouracil | ОН | Me | |
| acetyl | CH ₃ | H | H | 0 | 4-(N,N-diacetyl)cytosine | Н | Br | |
| acetyl | CH ₃ | H | H | S | 4-(N,N-diacetyl)cytosine | Н | Br | |
| acetyl | CH ₃ | OH | H | 0 | 4-(N,N-diacetyl)cytosine | H | Br | |
| acetyl | CH ₃ | OH | H | S | 4-(N,N-diacetyl)cytosine | H | Br | |

Example 2: Preparation of 2'-C-methylriboadenine

The title compound was prepared according to a published procedure (R.E. Harry-O'kuru, J.M. Smith, and M.S. Wolfe, "A short, flexible route toward 2'-C-branched ribonucleosides", *J.Org. Chem.* 1997, <u>62</u>, 1754-1759) (Scheme 8).

5

Scheme 8

(a) Dess-Martin periodinane; (b) MeMgBr / TiCl₄; (c) BzCl, DMAP, Et₃N; (d) bis(trimethylsilyl)acetamide, N⁶-benzoyl adenine, TMSOTf; (e) NH₃ / MeOH

In a similar manner, but using the appropriate sugar and pyrimidine or purine bases, the following nucleosides of Formula II are prepared.

wherein:

| R ¹ | \mathbb{R}^2 | R ³ | X¹ | X ² | Y |
|----------------|----------------|----------------|----|----------------|-----------------|
| Н | Н | H | H | H | Н |
| Н | H | Н | Н | Н | NH ₂ |
| Н | Н | Н | H | Н | NH-cyclopropyl |
| Н | H | Н | H | H | NH-methyl |
| Н | H | Н | H | H | NH-ethyl |
| Н | H | Н | H | Н | NH-acetyl |
| Н | H | Н | H | H | ОН |
| H | Н | H | H | H | OMe |
| Н | Н | Н | H | Н | OEt |
| Н | Н | Н | Н | Н | O-cyclopropyl |
| H | H | Н | H | Н | O-acetyl |
| H | Н | H | H | Н | SH |
| H | Н | Н | Н | Н | SMe |
| H | Н | Н | H | H | SEt |
| Н | Н | Н | H | Н | S-cyclopropyl |
| Н | Н | Н | H | Н | F |
| H | Н | Н | H | H | Cl |

| WO 01/90121 | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | $\overline{\mathbf{X}^2}$ | PCT/US01/16671 Y |
|---------------|----------------|----------------|----------------|---------------------------|---------------------|
| H | Н | H | - H | H | Br |
| H | H | H | H | H | I |
| | | | | | |
| monophosphate | Н | H | Н | H | NH ₂ |
| monophosphate | H | H | H | H | NH-acetyl |
| monophosphate | H | H | H | H | NH-cyclopropyl |
| monophosphate | Н | H | H | H | NH-methyl |
| monophosphate | H | H | Н | Н | NH-ethyl |
| monophosphate | Н | Н | H | H | ОН |
| monophosphate | Н | Н | Н | Н | O-acetyl |
| monophosphate | H | H | H | H | OMe |
| monophosphate | Н | Н | Н | H | OEt |
| monophosphate | Н | H | . Н | Н | O-cyclopropyl |
| monophosphate | Н | Н | H | Н | SH |
| monophosphate | Н | Н | Н | H | SMe |
| monophosphate | Н | H | H | H | SEt |
| monophosphate | Н | H | Н | Н | S-cyclopropyl |
| monophosphate | H | Н | H | H | F |
| monophosphate | H | Н | Н | Н | Cl |
| monophosphate | Н | Н | Н | Н | Br |
| monophosphate | H | H | H | Н | I |
| diphosphate | Н | H | H | H | NH ₂ |
| diphosphate | H | H | H | H | NH-acetyl |
| diphosphate | Н | H | H | H | NH-cyclopropyl |
| diphosphate | H | Н | H | H | NH-methyl |
| diphosphate | H | H | H . | Н | NH-ethyl |
| diphosphate | Н | H | Н | H | OH |
| diphosphate | Н | H | H | H | O-acetyl |
| diphosphate | Н | H | Н | H | OMe |
| diphosphate | Н | Н | H | H | OEt |
| diphosphate | H | Н | H | H | O-cyclopropyl |
| diphosphate | Н | H | H | H | SH |

| WO 01/90121 | R ² | \mathbb{R}^3 | \mathbf{X}^{1} | X^2 | CT/US01/16671 Y |
|----------------|----------------|----------------|------------------|-------|--------------------|
| | | | | | |
| diphosphate | H | H | H | H | SMe |
| diphosphate | H | H | H | H | SEt |
| diphosphate | H | Н | H | H | S-cyclopropyl |
| diphosphate | H | H | H | Н | F |
| diphosphate | H | Н | H | Н | Cl |
| diphosphate | Н | Н | H | Н | Br |
| diphosphate | Н | Н | Н | Н | I |
| triphosphate | Н | H | H | Н | NH ₂ |
| triphosphate | Н | H | H | H | NH-acetyl |
| triphosphate | Н | Н | H | H | NH-cyclopropyl |
| triphosphate | Н | H | H | H | NH-methyl |
| triphosphate | Н | Н | Н | H | NH-ethyl |
| triphosphate . | Н | Н | H | H | OH |
| triphosphate | Н | H | Н | H | OMe |
| triphosphate | Н | Н | H | H | OEt |
| triphosphate | Н | Н | H | H | O-cyclopropyl |
| triphosphate | Н | Н | H | H | O-acetyl |
| triphosphate | Н | Н | Н | H | SH |
| triphosphate | Н | Н | H | H | SMe |
| triphosphate | Н | Н | H | H | SEt |
| triphosphate | Н | H | H | H | S-cyclopropyl |
| triphosphate | Н | Н | H | H | F |
| triphosphate | Н | Н | H | H | Cl |
| triphosphate | Н | Н | H | H | Br |
| triphosphate | Н | H | H | H | I |
| monophosphate | monophosphate | monophosphate | Н | H | NH ₂ |
| monophosphate | monophosphate | monophosphate | H | H | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | Н | H | OH |
| monophosphate | monophosphate | monophosphate | H | H | F |
| monophosphate | monophosphate | monophosphate | H | H | Cl |
| diphosphate | diphosphate | diphosphate | H | H | NH ₂ |

| H H H H Cl H F H H Cl H Cl H H Cl H H H H Br H NH2 H H H Br H NH-cyclopropyl H H H H Br H OH H Br H OH H H H Br H OH H H H Br H OH H H H H NH-cyclopropyl H H H H H NH-cyclopropyl H H H H H NH Br H NH-cyclopropyl H H H H H NH Br H NH-cyclopropyl | R ¹ | \mathbb{R}^2 | R ³ | X ¹ | X^2 | Y Y |
|--|----------------|----------------|----------------|-----------------|-------|-----------------|
| diphosphate diphosphate diphosphate H H Cl diphosphate diphosphate diphosphate H H H Cl triphosphate H H Cl H H H H H F H NH-2 H H H H F H OH H H H F H F H OH H H H F H F H Cl H H H H F H F H Cl H H H H F H Cl H NH-2 H H H H Cl H NH-2 H H H Cl H NH-2 H H H Cl H F H OH H H H Cl H F H OH H H H Cl H F H OH H H H C CL H OH H H H H CL H OH H H H C CL H OH H H H H CL H F H OH H H H H CL H CL H H F H NH-2 H H H H CL H CL H NH-2 H H H H CL H CL H NH-2 H H H H CL H CL H NH-2 H H H H H CL H NH-2 H NH-2 H H H H H H H H NH-2 H NH-2 H H H H H H H NH-2 H NH-2 H H H H H H NH-2 H NH-2 H H H H H NH-2 H NH-2 H NH-2 H H H H H NH-2 H NH-2 H NH-2 H NH-2 H NH-2 H H H H NH-2 H H H H NH-2 H NH-2 H NH-2 H H H H NH-2 H NH-2 H NH-2 H H H H NH-2 H NH-2 H NH-2 H H H H NH-2 H NH-2 H NH-2 H H H H NH-2 H NH-3 H NH-3 H NH-3 H NH-3 H NH-3 H NH-4 H NH-4 H N | diphosphate | diphosphate | diphosphate | Н | Н | NH-cyclopropyl |
| diphosphate diphosphate triphosphate H H H F H OH H H H H H H Cl H H H H H H H H H H H H H H H H H H H | diphosphate | diphosphate | diphosphate | H | H | ОН |
| triphosphate H H H Cl H H H H H F H NH-cyclopropyl H H H H H F H OH H H H F H OH H H H F H F H Cl H H H H F H Cl H H H H Cl H NH-cyclopropyl H H H H H Cl H Cl H NH-cyclopropyl H H H H H Cl H Cl H F H Cl H H H H Cl H Cl H H H H Cl H F Cl H F Cl H H Cl H H H H B Cl H Cl H F Cl H H NH-cyclopropyl H H H H H B Cl H NH-cyclopropyl H H H H H B Cl H NH-cyclopropyl H H H H H NH-cyclopropyl | diphosphate | diphosphate | diphosphate | Н | H | F |
| triphosphate H H H Cl H H H H H F H NH2clopropyl H H H H H H F H Cl H H H H H H Cl H H F H Cl H H F H NH2clopropyl H H H H H Br H NH2clopropyl H H H H H Br H Cl H H H H H Br H Cl H H H H H NH2 H NH2clopropyl H H H H NH2 H NH2clopropyl | diphosphate | diphosphate | diphosphate | Н | H | Cl |
| triphosphate triphosphate triphosphate H H F triphosphate triphosphate triphosphate H H F triphosphate triphosphate H H CI H H H H CI H H H H H F H NH-cyclopropyl H H H H F H F H OH H H H H F H OH | triphosphate | triphosphate | triphosphate | H | H | NH ₂ |
| triphosphate triphosphate triphosphate H H F triphosphate triphosphate H H Cl H H H H H Cl H H H H F H NH-cyclopropyl H H H H F H OH H H H H F H OH H H H H F H F H H H H F H Cl H H H H H Cl H NH-cyclopropyl H H H H H H NH-cyclopropyl H H H H H NH-cyclopropyl H H H H H F H H H H H F H H H | triphosphate | triphosphate | triphosphate | Н | H | NH-cyclopropyl |
| triphosphate triphosphate H H Cl H H H F H NH2 H H H F H NH-cyclopropyl H H H F H OH H H H F H F H H H F H F H H H H F H F H H H H Cl H NH2 H H H H H OH H H H H H H H Cl H NH2 H NH2 H H H H H H NH2 H NH2 H NH2 H NH2 H NH2 H NH2 H H H H H H H H H H | triphosphate | triphosphate | triphosphate | Н | H | ОН |
| H H H H H F H NH2 H H H F H OH H H H F H OH H H H F H CI H NH-cyclopropyl H H H H H CI H NH2 H H H H CI H OH H H H H CI H CI H H H H CI H CI H OH H H H H CI H CI H CI H OH H H H H CI H CI H F H OH H H H H CI H F H CI H F H CI H F H CI H H CI H H CI H CI H CI H CI H CI | triphosphate | triphosphate | triphosphate | H | H | F |
| H H H H F H OH H H H F H OH H H H F H OH H H H F H CI H H H H H CI H NH-cyclopropyl H H H H H CI H OH H H H CI H OH H H H H CI H CI H H H H CI H CI H H H H CI H CI | triphosphate | triphosphate | triphosphate | H | H | Cl |
| H H H F H OH H H H F H F H H H F H F H H H H CI H NH2 H H H H OH | Н | Н | Н | F | H | NH ₂ |
| H H H H H F H Cl H H H H H Cl H NH2 H H H H Cl H NH-cyclopropyl H H H H H Cl H Cl H OH H H H H Cl H Cl H OH H H H H Cl H F H Cl H Cl H F H H H H Br H NH2 H NH-cyclopropyl H H H H H Br H NH-cyclopropyl H H H H H Br H NH-cyclopropyl H H H H H NH Br H OH H H H H H Br H OH H H H H H Br H OH H H H H H Br H F H H H H H Br H Cl H H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H OH | Н | H | Н | F | H | NH-cyclopropyl |
| H H H H H Cl H H H H H Cl H NH2 H H H H Cl H NH2 H H H H Cl H OH H H H H Cl H F H H H H Cl H F H H H H Cl H Cl H Cl H H H H H H Cl H Cl | H | Н | Н | F | H | OH |
| H H H H H CI H NH2 H H H H H CI H NH-cyclopropyl H H H H H H CI H F H H H H CI H CI H CI H CI H H CI H H CI H H CI H H NH-cyclopropyl H H H H H Br H NH-cyclopropyl H H H H H Br H CI H F H H H H H Br H CI H H F H H H H H Br H CI H H H H H Br H CI H H H H H Br H CI H H H H H H F H H H H H NH2 H NH2 H NH2 H NH-cyclopropyl H H H H H NH2 H NH-cyclopropyl H H H H H NH2 H NH-cyclopropyl H H H H H NH2 H OH | H | Н | H | F | H | F |
| H H H H CI H OH H H H CI H F H H H H CI H CI H H H H H CI H CI H H H H H Br H NH-cyclopropyl H H H H H Br H OH H H H H Br H OH H H H Br H OH H H H H Br H OH H H H H Br H OH H H H H Br H CI H H H H H Br H F H H H H H Br H CI H H H H H Br H CI H H H H H H Br H CI H H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H OH H H H NH2 H OH H H H NH2 H OH | Н | Н | Н | F | H | Cl |
| H H H H CI H F H H H H CI H CI H H H H CI H CI H H H H H H CI H NH2 H H H H H Br H NH-cyclopropyl H H H H H H Br H F H H H H H R Br H CI H H H H H NH2 H NH2 H H H H NH2 H NH-cyclopropyl H H H H H NH2 H NH2 H NH-cyclopropyl H H H H H NH2 H NH2 H F H H H H NH2 H OH | H | H | H | C1 | H | NH ₂ |
| H H H H H Cl H Cl H H H H H Br H NH2 H H H H Br H OH H H H Br H OH H H H H Br H Cl H H H H H Br H Cl H H H H H F H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H OH H NH2 H OH H H H NH2 H OH | H | H | H | Cl | H | NH-cyclopropyl |
| H H H H H CI H NH2 H H H H Br H NH-cyclopropyl H H H H H Br H OH H H H F H H H H F H H H H H NH2 H CI H H H H H H NH2 H OH H H H H H NH2 H NH2 H NH-cyclopropyl H NH2 H NH-cyclopropyl H NH2 H NH-cyclopropyl H H H H NH2 H NH2 H NH2 H OH H H H NH2 H OH H H H NH2 H OH | Н | H | Н | Cl | Н | ОН |
| H H H Br H NH2 H H H Br H NH-cyclopropyl H H H Br H OH H H H Br H F H H H Br H Cl H H H NH2 H NH2 H H H NH2 H OH H H H NH2 H F H H H NH2 H F H H H NH2 H Cl H H H NH2 H Cl H H H NH2 H NH2 | Н | H | Н | C1 | H | F |
| H H H Br H NH-cyclopropyl H H H Br H OH H H H Br H F H H H Br H Cl H H H NH2 H NH2 H H H NH2 H NH-cyclopropyl H H H NH2 H OH H H H NH2 H F H H H NH2 H Cl H H H NH2 H Cl H H H NH2 H NH2 | Н | H | Н | C1 | Н | Cl |
| H H H H H Br H F H H H H Br H Cl H H H H NH2 H NH2 H NH-cyclopropyl H H H H NH2 H OH H NH2 H Cl H NH2 H Cl H NH2 H Cl H NH2 H NH-cyclopropyl H H H H NH2 H OH H NH2 H Cl H H H NH2 H OH | H | Н | Н | Br | H | NH ₂ |
| H H H H H Br H F H H H H NH ₂ H NH-cyclopropyl H H H H NH ₂ H OH H H H NH ₂ H OH H H H NH ₂ H OH H H H NH ₂ H CI H H H H NH ₂ H OH H H H NH ₂ H F | Н | Ĥ | Н | Br | H | NH-cyclopropyl |
| H H H H H NH ₂ H NH-cyclopropyl H H H H NH ₂ H OH H NH-cyclopropyl H H H H NH ₂ H F H H H NH ₂ H Cl H H H NH ₂ H F | Н | H | H | Br | H | ОН |
| H H H H NH ₂ H NH ₂ H NH-cyclopropyl H H H H NH ₂ H OH H H H NH ₂ H CI H H H H NH ₂ H NH-cyclopropyl | H | H | Н | Br | H | F |
| H H H NH2 H NH-cyclopropyl H H H NH2 H OH H H H NH2 H F H H H NH2 H Cl H H H SH H NH2 | H | H | Н | Br | Н | Cl |
| H H H NH2 H OH H H H NH2 H F H H H NH2 H Cl H H H SH H NH2 | H | H | Н | NH ₂ | Н | NH ₂ |
| H H H NH ₂ H F H H H NH ₂ H Cl H H H SH H NH ₂ | H | Н | Н | NH ₂ | Н | NH-cyclopropyl |
| H H H NH ₂ H Cl H H H SH H NH ₂ | Н | H | Н | NH ₂ | H | ОН |
| H H H SH H NH2 | H | H | Н | NH ₂ | Н | F |
| | H | H | Н | NH ₂ | Н | Cl |
| H H SH H NH-cyclopropyl | H | H | Н | SH | H | NH ₂ |
| | H | H | Н | SH | Н | NH-cyclopropyl |

| \mathbb{R}^1 | R ² | \mathbb{R}^3 | X ¹ | X^2 | Y |
|----------------|----------------|----------------|----------------|-------|-----------------|
| Н | Н | H | SH | H | ОН |
| Н | Н | Н | SH | Н | F |
| Н | H | H | SH | H | Cl |
| acetyl | H | H | H | H | NH ₂ |
| acetyl | H | H | H | H | NH-cyclopropyl |
| acetyl | Н | H | H | H | OH |
| acetyl | Н | H | H | H | F |
| acetyl | Н | Н | H | H | Cl |
| acetyl | Н | H | F | H | NH ₂ |
| acetyl | H | Н | F | H | NH-cyclopropyl |
| acetyl | H | H | F | H | OH |
| acetyl | Н | Н | F | H | F |
| acetyl | H | Н | F | H | Cl |
| H | acetyl | acetyl | H | H | NH ₂ |
| Н | acetyl | acetyl | H | H | NH-cyclopropyl |
| Н | acetyl | acetyl | H | H | OH |
| Н | acetyl | acetyl | H | H | F |
| Н | acetyl | acetyl | H | H | Cl |
| acetyl | acetyl | acetyl | H | H | NH ₂ |
| acetyl | acetyl | acetyl | H | Н | NH-cyclopropyl |
| acetyl | acetyl | acetyl | H | H | OH |
| acetyl | acetyl | acetyl | H | H | F |
| acetyl | acetyl | acetyl | Н | H | Cl |
| monophosphate | acetyl | acetyl | Н | H | NH ₂ |
| monophosphate | acetyl | acetyl | H | Н | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | Н | Н | OH |
| monophosphate | acetyl | acetyl | Н | H | F |
| monophosphate | acetyl | acetyl | H | H | Cl |
| diphosphate | acetyl | acetyl | H | H | NH ₂ |
| diphosphate | acetyl | acetyl | Н | H | NH-cyclopropy |
| diphosphate | acetyl | acetyl | Н | H | OH |

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|----------------|----------------|----------------|-----|-----------------|-----------------|
| \mathbb{R}^1 | R ² | R ³ | X1 | X ² | Y |
| diphosphate | acetyl | acetyl | H | H | F |
| diphosphate | acetyl | acetyl | H | H | Cl |
| triphosphate | acetyl | acetyl | Н | H | NH ₂ |
| triphosphate | acetyl | acetyl | H | H | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | H | H | ОН |
| triphosphate | acetyl | acetyl | Н | H | F |
| triphosphate | acetyl | acetyl | H | Н | Cl |
| Н | Н | H | Н | NH ₂ | Н |
| Н | H | H | Н | NH ₂ | NH ₂ |
| H | Н | Н | Н | NH ₂ | NH-cyclopropyl |
| H | Н | Н | Н | NH ₂ | NH-methyl |
| Н | H | H | Н | NH ₂ | NH-ethyl |
| Н | Н | Н | H | NH ₂ | NH-acetyl |
| Н | Н | Н | H | NH ₂ | OH |
| H | Н | H | H | NH ₂ | OMe |
| Н | Н | Н | H | NH ₂ | OEt |
| Н | Н | H | H | NH ₂ | O-cyclopropyl |
| Н | Н | Н | Н | NH ₂ | O-acetyl |
| Н | H | Н | H | NH ₂ | SH |
| Н . | Н | H | Н | NH ₂ | SMe |
| H | Н | H | H | NH ₂ | SEt |
| H | Н | Н | H | NH ₂ | S-cyclopropyl |
| Н | Н | Н | Н | NH ₂ | F |
| H | Н | Н | H | NH ₂ | Cl |
| H | H | Н | H | NH ₂ | Br |
| H | H | Н | Н | NH ₂ | I |
| monophosphate | H | Н | H | NH ₂ | NH ₂ |
| monophosphate | H | Н | Н | NH ₂ | NH-acetyl |
| monophosphate | H | H | · H | NH ₂ | NH-cyclopropyl |
| monophosphate | H | H | H | NH ₂ | NH-methyl |
| monophosphate | Н | H | H | NH ₂ | NH-ethyl |

WO 01/90121 PCT/US01/16671 $\overline{X^1}$ $\overline{X^2}$ R^2 $\overline{\mathbf{R}^3}$ \mathbf{R}^{1} H H NH₂ OH H monophosphate H H O-acetyl NH_2 monophosphate H NH₂ H Η OMe monophosphate H H H H NH₂ OEt monophosphate NH₂ Η H H O-cyclopropyl monophosphate H H NH_2 SH monophosphate H H Н H $\overline{\mathrm{NH_2}}$ SMe monophosphate H H NH₂ SEt monophosphate Η S-cyclopropyl NH₂ H H Н monophosphate H Η H NH₂ monophosphate H H Cl H NH_2 monophosphate H H Η NH_2 \mathbf{Br} monophosphate $\overline{\mathbf{H}}$ Н NH_2 Ī H monophosphate Η H Η NH₂ NH_2 diphosphate H NH₂ NH-acetyl diphosphate Н Н diphosphate H H Н NH₂ NH-cyclopropyl H NH_2 NH-methyl H diphosphate Η H Η NH_2 NH-ethyl diphosphate H H NH₂ OH Н Н diphosphate H $\overline{\mathbf{H}}$ NH₂ O-acetyl diphosphate Η H NH₂ OMe diphosphate Η H

H NH₂ OEt diphosphate H H H H NH₂ O-cyclopropyl diphosphate Η H Н NH_2 SH diphosphate Н SMe H Η NH_2 diphosphate Η H H NH₂ SEt diphosphate H H Н NH_2 S-cyclopropyl diphosphate Η Н H H NH_2 F diphosphate C1 Η H H NH_2 diphosphate Н H H NH₂ Br diphosphate H NH₂ Η Ι diphosphate Η

| WO 01/90121 | T=2 | T-3 | 771 | | CT/US01/16671 |
|----------------|----------------|----------------|----------------|-----------------|-----------------|
| R ¹ | R ² | R ³ | X ¹ | X ² | Y |
| triphosphate | H | Н | H | NH ₂ | NH ₂ |
| triphosphate | Н | Н . | H | NH ₂ | NH-acetyl |
| triphosphate | Н | H | H | NH ₂ | NH-cyclopropyl |
| triphosphate | Н | Н | H | NH ₂ | NH-methyl |
| triphosphate | H · | H | H | NH ₂ | NH-ethyl |
| triphosphate | Н | Н | H | NH ₂ | ОН |
| triphosphate | Н | Н | H | NH ₂ | OMe |
| triphosphate | Н | H | H | NH ₂ | OEt |
| triphosphate | Н | Н | H | NH ₂ | O-cyclopropyl |
| triphosphate | H | H | H | NH ₂ | O-acetyl |
| triphosphate | Н | H | H | NH ₂ | SH |
| triphosphate | Н | H | H | NH ₂ | SMe |
| triphosphate | H | H | H | NH ₂ | SEt |
| triphosphate | H | Н | H | NH ₂ | S-cyclopropyl |
| triphosphate | Н | Н | H | NH ₂ | F |
| triphosphate | Н | Н | H | NH ₂ | Cl |
| triphosphate | Н | H | H | NH ₂ | Br |
| triphosphate | Н | Н | H | NH ₂ | I |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | NH ₂ |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | ОН |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | F |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | Cl |
| diphosphate | diphosphate | diphosphate | Н | NH ₂ | NH ₂ |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | ОН |
| diphosphate | diphosphate | diphosphate | Н | NH ₂ | F |
| diphosphate | diphosphate | diphosphate | Н | NH ₂ | Cl |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | NH ₂ |
| triphosphate | triphosphate | triphosphate | Н | NH ₂ | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | Н | NH ₂ | ОН |
| | | | | | |

| R ¹ | \mathbb{R}^2 | | 1 V - | 1 V- | Y |
|----------------|----------------|----------------|-----------------|-----------------|-----------------|
| | | R ³ | X ¹ | X ² | |
| triphosphate | triphosphate | triphosphate | Н | NH ₂ | F |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | Cl |
| H | H | Н | F | NH ₂ | NH ₂ |
| Н | Н | Н | F | NH ₂ | NH-cyclopropyl |
| Н | Н | Н | F | NH ₂ | OH |
| Н | Н | Н | F | NH ₂ | F |
| Н | Н | Н | F | NH ₂ | Cl |
| Н | Н | Н | Cl | NH ₂ | NH ₂ |
| Н | Н | Н | Cl | NH ₂ | NH-cyclopropyl |
| H | Н | Н | Cl | NH ₂ | ОН |
| H | Н | Н | Cl | NH ₂ | F |
| H | Н | Н | C1 | NH ₂ | Cl |
| Н | Н | Н | Br | NH ₂ | NH ₂ |
| H | Н | Н | Br | NH ₂ | NH-cyclopropyl |
| Н | Н | Н | Br | NH ₂ | ОН |
| Н | Н | Н | Br | NH ₂ | F |
| Н | H | Н | Br | NH ₂ | Cl |
| Н | Н | Н | NH ₂ | NH ₂ | NH ₂ |
| H | Н | Н | NH ₂ | NH ₂ | NH-cyclopropyl |
| H | Н | Н | NH ₂ | NH ₂ | ОН |
| Н | H | Н | NH ₂ | NH ₂ | F |
| Н | H | Н | NH ₂ | NH ₂ | Cl |
| H | Н | H | SH | NH ₂ | NH ₂ |
| H | Н | Н | SH | NH ₂ | NH-cyclopropyl |
| H | H | H . | SH | NH ₂ | ОН |
| H | H | H | SH | NH ₂ | F |
| H | Н | H | SH | NH ₂ | Cl |
| acetyl | Н | H | Н | NH ₂ | NH ₂ |
| acetyl | Н | Н | H | NH ₂ | NH-cyclopropyl |
| acetyl | H | Н | H | NH ₂ | ОН |
| acetyl | H | Н | H | NH ₂ | F |

| \mathbb{R}^1 | R ² | \mathbb{R}^3 | X ¹ | X ² | $\overline{\mathbf{Y}}^-$ |
|----------------|----------------|----------------|----------------|-----------------|---------------------------|
| acetyl | Н | H | H | NH ₂ | Cl |
| acetyl | H | Н | F | NH ₂ | NH ₂ |
| acetyl | H | H | F | NH ₂ | NH-cyclopropyl |
| acetyl | H | Н | F | NH ₂ | OH |
| acetyl | H | Н | F | NH ₂ | F |
| acetyl | Н | Н | F | NH ₂ | Cl |
| Н | acetyl | acetyl | H | NH ₂ | NH ₂ |
| Н | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| Н | acetyl | acetyl | H | NH ₂ | OH |
| Н | acetyl | acetyl | H | NH ₂ | F |
| Н | acetyl | acetyl | Н | NH ₂ | Cl · |
| acetyl | acetyl | acetyl | Н | NH ₂ | NH ₂ |
| acetyl | acetyl | acetyl | Н | NH ₂ | NH-cyclopropyl |
| acetyl | acetyl | acetyl | Н | NH ₂ | OH |
| acetyl | acetyl | acetyl | Н | NH ₂ | F |
| acetyl | acetyl | acetyl | Н | NH ₂ | Cl |
| monophosphate | acetyl | acetyl | Н | NH ₂ | NH ₂ |
| monophosphate | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | Н | NH ₂ | ОН |
| monophosphate | acetyl | acetyl | Н | NH ₂ | F |
| monophosphate | acetyl | acetyl | Н | NH ₂ | Cl |
| diphosphate | acetyl | acetyl | H | NH ₂ | NH ₂ |
| diphosphate | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | H | NH ₂ | ОН |
| diphosphate | acetyl | acetyl | Н | NH ₂ | F |
| diphosphate | acetyl | acetyl | Н | NH ₂ | Cl |
| triphosphate | acetyl | acetyl | Н | NH ₂ | NH ₂ |
| triphosphate | acetyl | acetyl | H | NH ₂ | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | H | NH ₂ | OH |
| triphosphate | acetyl | acetyl | H | NH ₂ | F |
| triphosphate | acetyl | acetyl | H | NH ₂ | Cl |

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|----------------|----------------|----------------|----|----------------|-----------------|
| R ¹ | R ² | R ³ | X1 | X ² | Y |
| Н | Н | Н | H | Cl | Н |
| Н | Н | Н | H | Cl | Н |
| Н | Н | Н | H | Cl | NH ₂ |
| Н | Н | Н | H | Cl | NH-cyclopropyl |
| Н | Н | Н | H | C1 | NH-methyl |
| Н | Н | Н | H | Cl | NH-ethyl |
| H | Н | Н | H | Cl | NH-acetyl |
| H | H | Н | H | Cl | ОН |
| H | Н | H | H | Cl | OMe |
| H | Н | Н | H | Cl | OEt |
| Н | Н | H | Н | Cl | O-cyclopropyl |
| H | Н | H | Н | Cl | O-acetyl |
| H | Н | Н | H | C1 | SH |
| Н | Н | H | H | C1 | SMe |
| H | Н | Н | H | Cl | SEt |
| H | Н | Н | H | Cl | S-cyclopropyl |
| monophosphate | Н | Н | H | C1 | NH ₂ |
| monophosphate | Н | H | H | C1 | NH-acetyl |
| monophosphate | Н | Н | H | Cl | NH-cyclopropyl |
| monophosphate | Н | Н | H | C1 | NH-methyl |
| monophosphate | H | Н | H | Cl | NH-ethyl |
| monophosphate | H | H | H | Cl | ОН |
| monophosphate | Н | H | H | Cl | O-acetyl |
| monophosphate | Н | Н | H | Cl | OMe |
| monophosphate | Н | H | H | Cl | OEt |
| monophosphate | Н | Н | H | Cl | O-cyclopropyl |
| monophosphate | Н | Н | H | Cl | SH |
| monophosphate | Н | Н | H | Cl | SMe |
| monophosphate | Н | Н | H | Cl | SEt |
| monophosphate | Н | H | H | Cl | S-cyclopropyl |
| diphosphate | Н | Н | H | Cl | NH ₂ |

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|----------------|----------------|----------------|----------------|----------------|-----------------|
| \mathbb{R}^1 | R ² | R ³ | X ¹ | X ² | Y |
| diphosphate | Н | Н | Н | C1 | NH-acetyl |
| diphosphate | Н | Н | H | Cl | NH-cyclopropyl |
| diphosphate | Н | Н | Н | Cl | NH-methyl |
| diphosphate | Н | Н | Н | Cl | NH-ethyl |
| diphosphate | H | Н | H | Cl | ОН |
| diphosphate | Н | Н | H | Cl | O-acetyl |
| diphosphate | Н | Н | H | Cl | ОМе |
| diphosphate | Н | Н | H | Cl | OEt |
| diphosphate | Н | Н | H | Cl | O-cyclopropyl |
| diphosphate | Н | Н | H | Cl | SH |
| diphosphate | Н | Н | H | Cl | SMe |
| diphosphate | Н | Н | H | Cl | SEt |
| diphosphate | Н | H | H | Cl | S-cyclopropyl |
| triphosphate | H | Н | H | Cl | NH ₂ |
| triphosphate | H | Н | H | Cl | NH-acetyl |
| triphosphate | Н | Н | H | Cl | NH-cyclopropyl |
| triphosphate | Н | Н | H | Cl | NH-methyl |
| triphosphate | Н | H | Н | Cl | NH-ethyl |
| triphosphate | H | Н | H | Cl | ОН |
| triphosphate | H | H | Н | Cl | OMe |
| triphosphate | Н | Н | H | Cl | OEt |
| triphosphate | H | Н | H | Cl | O-cyclopropyl |
| triphosphate | H | H | H | Cl | O-acetyl |
| triphosphate | Н | Н | H | Cl | SH |
| triphosphate | H | Н | H | Cl | SMe |
| triphosphate | H | Н . | Н | Cl | SEt |
| triphosphate | Н | H | H | C1 | S-cyclopropyl |
| monophosphate | monophosphate | monophosphate | Н | Cl | NH ₂ |
| monophosphate | monophosphate | monophosphate | Н | Cl | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | Cl | OH |
| diphosphate | diphosphate | diphosphate | H | Cl | NH ₂ |

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|----------------------------|----------------|----------------|-----------------|----------------|-----------------|
| R ¹ | R ² | R ³ | X ¹ | X ² | Y |
| diphosphate | diphosphate | diphosphate | H | CI | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | Cl | ОН |
| triphosphate | triphosphate | triphosphate | H | Cl | NH ₂ |
| triphosphate | triphosphate | triphosphate | Н | Cl | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | Н | Cl | ОН |
| H | Н | Н | F | Cl | NH ₂ |
| Н | Н | Н | F | Cl | NH-cyclopropyl |
| Н | Н | Н | F | Cl | ОН |
| Н | Н | Н | Cl | Cl | NH ₂ |
| Н | Н | Н | C1 | Cl | NH-cyclopropyl |
| Н | Н | H | Cl | Cl | ОН |
| Н | Н | Н | Br | Cl | NH ₂ |
| Н | H | Н | Br | Cl | NH-cyclopropyl |
| Н | Н | Н | Br | Cl | ОН |
| H | Н | Н | NH ₂ | Cl | NH ₂ |
| Н | Н | Н | NH ₂ | Cl | NH-cyclopropyl |
| H | Н | Н | NH ₂ | Cl | ОН |
| H | Н | Н | SH | Cl | NH ₂ |
| Н | Н | Н | SH | C1 | NH-cyclopropyl |
| H | Н | Н | SH | C1 | ОН |
| acetyl | Н | Н | H | Cl | NH ₂ |
| acetyl | Н | Н | H | Cl | NH-cyclopropyl |
| acetyl | Н | Н | Н | Cl | ОН |
| acetyl | Н | H | F | Cl | NH ₂ |
| acetyl | H | H | F | Cl | NH-cyclopropyl |
| acetyl | Н | H | F | Cl | ОН |
| Н | acetyl | acetyl | H | Cl | NH ₂ |
| Н | acetyl | acetyl | H | Cl | NH-cyclopropyl |
| H | acetyl | acetyl | H | Cl | ОН |
| acetyl | acetyl | acetyl | H | Cl | NH ₂ |
| acetyl | acetyl | acetyl | Н | Cl | NH-cyclopropyl |
| Ļ | | | | | |

WO 01/90121 PCT/US01/16671 \mathbb{R}^3 $\overline{X^1}$ $\overline{X^2}$ $\overline{\mathbf{R^2}}$ R^{T} Y acetyl H Cl OH acetyl acetyl H Cl NH_2 monophosphate acetyl acetyl H Cl NH-cyclopropyl acetyl monophosphate acetyl H Cl OH monophosphate acetyl acetyl Cl NH₂ H diphosphate acetyl acetyl Cl Η NH-cyclopropyl diphosphate acetyl acetyl H Cl OH diphosphate acetyl acetyl H Cl NH₂ triphosphate acetyl acetyl Cl H NH-cyclopropyl triphosphate acetyl acetyl OH Η Cl triphosphate acetyl acetyl H H H Cl NH_2 Η Cl H H NH-cyclopropyl H Η OH H Н Н C1 Н NH₂ Н Н H Br Η

Alternatively, the following nucleosides of Formula V are prepared, using the appropriate sugar and pyrimidine or purine bases.

Η

H

H

Η

H

Н

Br

Br

NH-cyclopropyl

OH

$$X^1$$
 X^1
 X^1

wherein:

5

H

Н

| R ¹ | \mathbb{R}^2 | R ³ | X ¹ | Y |
|----------------|----------------|----------------|----------------|-----------------|
| H | H | H | H | Н |
| H | Н . | Н | H | NH ₂ |

 $\overline{X^1}$ $\overline{\mathbf{R^2}}$ R^3 \mathbb{R}^1 . Y H H H H NH-cyclopropyl Н H H H NH-methyl H H H H NH-ethyl Н H NH-acetyl H H H H OH H H H H H H OMe H OEt $\overline{\mathbf{H}}$ \overline{H} Η H H O-cyclopropyl H H Н H O-acetyl H H SH H H Η H H SMe Н Н H H \overline{H} SEt H Η H H Н H S-cyclopropyl Н H NH₂ Η monophosphate H NH-acetyl H H monophosphate H NH-cyclopropyl monophosphate H H Н NH-methyl Н H monophosphate Н NH-ethyl H H monophosphate \overline{H} Н OH H monophosphate Н O-acetyl Н H monophosphate H OMe H H monophosphate H OEt Н H monophosphate H O-cyclopropyl H H monophosphate H H SH monophosphate H Н H SMe H monophosphate H H H SEt monophosphate $\overline{\mathbf{H}}$ H S-cyclopropyl H monophosphate diphosphate Н H H NH_2 Н NH-acetyl Η Н diphosphate H H NH-cyclopropyl diphosphate Н H NH-methyl diphosphate Н H

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| R ¹ | \mathbb{R}^2 | R ³ | X | PCT/US01/166 Y |
|----------------|----------------|----------------|---|-------------------|
| diphosphate | H | H | Н | NH-ethyl |
| diphosphate | H | H | H | OH |
| diphosphate | H | H | Н | O-acetyl |
| diphosphate | Н | H | Н | OMe |
| diphosphate | H | H | H | OEt |
| diphosphate | H | H | Н | O-cyclopropyl |
| diphosphate | H | H | Н | SH |
| diphosphate | Н | H | Н | SMe |
| diphosphate | Н | H | H | SEt |
| diphosphate | H | H | H | S-cyclopropyl |
| triphosphate | H | Н | H | NH ₂ |
| triphosphate | H | H | Н | NH-acetyl |
| triphosphate | Н | H | Н | NH-cyclopropyl |
| triphosphate | H | Н | H | NH-methyl |
| triphosphate | H | H | H | NH-ethyl |
| triphosphate | H | Н | H | ОН |
| triphosphate | Н | Н | Н | OMe |
| triphosphate | H | H | H | OEt |
| triphosphate | Н | Н | H | O-cyclopropyl |
| triphosphate | Н | H | Н | O-acetyl |
| triphosphate | Н | Н | H | SH |
| triphosphate | Н | H | H | SMe |
| triphosphate | Н | Н | Н | SEt |
| triphosphate | Н | Н | H | S-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | NH ₂ |
| monophosphate | monophosphate | monophosphate | Н | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | OH |
| diphosphate | diphosphate | diphosphate | Н | NH ₂ |
| diphosphate | diphosphate | diphosphate | H | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | OH |
| triphosphate | triphosphate | triphosphate | H | NH ₂ |
| | L | | | · |

| R ¹ | R ² | R ³ | X¹ | PCT/US01/166 |
|----------------|----------------|----------------|-----------------|-----------------|
| triphosphate | triphosphate | triphosphate | H | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | Н | ОН |
| Н | Н | Н | F | NH ₂ |
| H | Н | Н | F | NH-cyclopropyl |
| Н | Н | Н | F | ОН |
| Н | H | Н | Cl | NH ₂ |
| Н | H | Н | Cl | NH-cyclopropyl |
| Н | H | H | Cl | ОН |
| Н | H | Н | Br | NH ₂ |
| Н | H | Н | Br | NH-cyclopropyl |
| Н | H | Н | Br | ОН |
| Н | H . | H | NH ₂ | NH ₂ |
| Н | Н | H | NH ₂ | NH-cyclopropyl |
| H | Н | Н | NH ₂ | OH |
| H | Н | Н | SH | NH ₂ |
| Н | Н | Н | SH | NH-cyclopropyl |
| H | Н | Н | SH | OH |
| acetyl | Н | Н | H | NH ₂ |
| acetyl | Н | Н | Н | NH-cyclopropyl |
| acetyl | Н | Н | H | ОН |
| acetyl | H | Н | F | NH ₂ |
| acetyl | H | Н | F | NH-cyclopropyl |
| acetyl | H | Н | F | OH |
| Н | acetyl | acetyl | H | NH ₂ |
| Н | acetyl | acetyl | H | NH-cyclopropyl |
| Н | acetyl | acetyl | H | OH |
| acetyl | acetyl | acetyl | Н | NH ₂ |
| acetyl | acetyl | acetyl | Н | NH-cyclopropyl |
| acetyl | acetyl | acetyl | H | ОН |
| monophosphate | acetyl | acetyl | Н | NH ₂ |
| monophosphate | acetyl | acetyl | H | NH-cyclopropyl |

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|------------------|----------------|----------------|----------------|-----------------|
| \mathbb{R}^{1} | R ² | R ³ | X ¹ | Y |
| monophosphate | acetyl | acetyl | Н | OH |
| diphosphate | acetyl | acetyl | H | NH ₂ |
| diphosphate | acetyl | acetyl | Н | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | Н | ОН |
| triphosphate | acetyl | acetyl | H | NH ₂ |
| triphosphate | acetyl | acetyl | H | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | Н | OH |

Alternatively, the following nucleosides of Formula X are prepared, using the appropriate sugar and pyrimidine or purine bases.

$$R^{1}O$$
 X
 R^{6}
 OR^{2}
 OR^{3}
 (X)

wherein:

| R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | R ⁶ | · X | Base |
|----------------|----------------|----------------|-----------------|-----|-------------------|
| H | H | Н | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetyluracil |
| Н | Н . | Н | CH ₃ | 0 | Hypoxanthine |
| Н | H | H | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthymine |
| H | Н | Н | CH ₃ | 0 | Thymine |
| H | Н | H | CH ₃ | 0 | Cytosine |
| H | H | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| H | H | H | CH ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| H | H | Н | CH ₃ | 0 | Uracil |

| H H H CH ₃ S Thymin H H H CH ₃ S Cytosin H H H CH ₃ S 4-(N-m acetyl) acetyl) H H H CH ₃ S 4-(N,N diacety H H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | 1/16671 |
|--|------------|
| H H H H CH3 S 2,4-O-Diacety H H H H CH3 S Hypoxa H H H H CH3 S 2,4-O-Diacety H H H H H CH3 S Thymin H H H CH3 S Cytosin H H H H CH3 S Cytosin H H H H CH3 S 4-(N-m acetyl) H H H H H CH3 S Uracil H H H H CH3 S Uracil H H H H CH3 S 5-Fluor monophosphate H H CH3 O 2,4-O- | rouracil |
| H H H CH3 S Hypox H H H CH3 S 2,4-O-Diacety H H H CH3 S Thymin H H H CH3 S Cytosin H H H CH3 S 4-(N-macetyl) H H H CH3 S 4-(N,N) diacetyl H H CH3 S Uracil H H H CH3 S 5-Fluor monophosphate H H CH3 O 2,4-O- | ···· |
| H H H CH3 S Hypox H H H CH3 S 2,4-O-Diacety H H H CH3 S Thymin H H H CH3 S Cytosin H H H CH3 S 4-(N-max) acetylo A CH3 S 4-(N,N) H H H CH3 S Uracil H H H CH3 S 5-Fluor monophosphate H H CH3 O 2,4-O- | |
| H H H CH ₃ S 2,4-O-Diacety H H H CH ₃ S Thymin H H H CH ₃ S Cytosin H H H CH ₃ S 4-(N-m acetyl) A CH ₃ S 4-(N,N diacety B H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | |
| H H H CH ₃ S Thymin H H H CH ₃ S Cytosin H H H CH ₃ S 4-(N-m H H H CH ₃ S 4-(N,N diacety H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | anthine |
| H H H CH ₃ S Thymin H H H CH ₃ S Cytosin H H H CH ₃ S 4-(N-m acetyl) acetyl) H H H CH ₃ S 4-(N,N diacety H H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | |
| H H H CH ₃ S Cytosin H H H CH ₃ S 4-(N-m acetyl) H H CH ₃ S 4-(N,N) diacety H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | ylthymine |
| H H H CH ₃ S 4-(N-m acetyl)e H H H CH ₃ S 4-(N,N diacetyl)e H H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | ne |
| H H H H CH3 S 4-(N,N) H H H CH3 S Uracil H H H CH3 S 5-Fluor monophosphate H H CH3 O 2,4-O- | ne |
| H H H CH ₃ S 4-(N,N) diacety H H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | iono- |
| H H H CH3 S Uracil H H H CH3 S 5-Fluor monophosphate H H CH3 O 2,4-O- | cytosine |
| H H H CH ₃ S Uracil H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | - |
| H H H CH ₃ S 5-Fluor monophosphate H H CH ₃ O 2,4-O- | d)cytosine |
| monophosphate H H CH ₃ O 2,4-O- | |
| | rouracil |
| Diacet | |
| | yluracil |
| monophosphate H H CH ₃ O Hypox | anthine |
| monophosphate H H CH ₃ O 2,4-O- | |
| Diacet | ylthym |
| monophosphate H H CH ₃ O Thymir | ne |
| monophosphate H H CH ₃ O Cytosia | ne |
| monophosphate H H CH ₃ O 4-(N-m | iono- |
| acetyl) | cytosine |
| monophosphate H H CH ₃ O 4-(N,N | [- |
| diacety | l)cytosine |
| monophosphate H H CH ₃ O Uracil | ·_ |
| monophosphate H H CH ₃ O 5-Fluo | rouracil |
| monophosphate H H CH ₃ S 2,4-O- | |
| Diacet | yluracil |
| monophosphate H H CH ₃ S Hypox | anthine |
| monophosphate H H CH ₃ S 2,4-O- | anamic |
| Diacet | |

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|----------------|----------------|----------------|-----------------|---|-------------------|
| R ¹ | R ² | \mathbb{R}^3 | R ⁶ | X | Base |
| monophosphate | Н | H | CH ₃ | S | Thymine |
| monophosphate | Н | Н | CH ₃ | S | Cytosine |
| monophosphate | Н | H | CH ₃ | S | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| monophosphate | H | Н | CH ₃ | S | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| monophosphate | Н | Н | CH ₃ | S | Uracil |
| monophosphate | Н | Н | CH ₃ | S | 5-Fluorouracil |
| diphosphate | Н | Н | CH ₃ | 0 | 2,4-0- |
| | | | | | Diacetyluracil |
| diphosphate | H | H | CH ₃ | 0 | Hypoxanthine |
| diphosphate | Н | Н | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthymine |
| diphosphate | Н | H | CH ₃ | 0 | Thymine |
| diphosphate | Н | Н | CH ₃ | 0 | Cytosine |
| diphosphate | Н | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | } | acetyl)cytosine |
| diphosphate | Н | H | CH ₃ | 0 | 4-(N,N- |
| | | | | į | diacetyl)cytosine |
| diphosphate | Н | H | CH ₃ | 0 | Uracil |
| diphosphate | Н | H | CH ₃ | 0 | 5-Fluorouracil |
| diphosphate | Н | Н | CH ₃ | S | 2,4-O- |
| · | | | | | Diacetyluracil |
| diphosphate | Н | H | CH ₃ | S | Hypoxanthine |
| diphosphate | H | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetylthym |
| diphosphate | H | Н | CH ₃ | S | Thymine |
| diphosphate | Н | H . | CH ₃ | S | Cytosine |
| triphosphate | Н | Н | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetyluracil |
| triphosphate | Н | Н | CH ₃ | 0 | Hypoxanthine |

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|----------------|----------------|----------------|-----------------|---|-------------------|
| \mathbb{R}^1 | R ² | \mathbb{R}^3 | R ⁶ | X | Base |
| triphosphate | Н | H | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthymine |
| triphosphate | Н | Н | CH ₃ | 0 | Thymine |
| triphosphate | H | Н | CH ₃ | 0 | Cytosine |
| triphosphate | Н | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| triphosphate | Н | H | CH ₃ | 0 | 4-(N,N- |
| | | | : | | diacetyl)cytosine |
| triphosphate | Н | Н | CH ₃ | 0 | Uracil |
| triphosphate | Н | Н | CH ₃ | 0 | 5-Fluorouracil |
| triphosphate | Н | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetyluracil |
| triphosphate | Н | H | CH ₃ | S | Hypoxanthine |
| triphosphate | Н | H | CH ₃ | S | 2,4-O- |
| | | | | | Diacetylthymine |
| triphosphate | Н | H | CH ₃ | S | Thymine |
| triphosphate | Н | H · | CH ₃ | S | Cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,4-O- |
| | | | | | Diacetyluracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Hypoxanthine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Thymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | Uracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 5-Fluorouracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,4-0- |
| | | | | | Diacetyluracil |

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|----------------|----------------|----------------|-----------------|---|-------------------|
| R ¹ | R ² | R ³ | R ⁶ | X | Base |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | Hypoxanthine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,4-O- |
| | | | | | Diacetylthymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | Thymine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | Cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 4-(N-mono- |
| | | | _ | | acetyl)cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | Uracil |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 5-Fluorouracil |
| acetyl | acetyl | acetyl | CF ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| acetyl | acetyl | acetyl | CF ₃ | S | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| acetyl | acetyl | acetyl | 2-bromo- | 0 | 4-(N,N- |
| | | | vinyl | | diacetyl)cytosine |
| acetyl | acetyl | acetyl | 2-bromo- | S | 4-(N,N- |
| | | | vinyl | | diacetyl)cytosine |
| H | Н | Н | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| H | H | H | CH ₃ | 0 | 6-O-acetyl |
| | | | | | guanine |
| Н | H | Н | CH ₃ | 0 | 8-fluoroguanine |
| Н | Н | H | CH ₃ | 0 | guanine |
| H | Н | Н | CH ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| Н | Н | H | CH ₃ | 0 | 2-fluoroadenine |
| Н | Н | Н | CH ₃ | 0 | 8-fluoroadenine |
| H | H | H | CH ₃ | 0 | 2,8-difluoro- |
| | | | | | adenine |
| Н | H | H | CH ₃ | 0 | adenine |

| | | | | PCT/US01/16671 | |
|----------------|----------------|----------------|-----------------|----------------|-------------------|
| R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | R ⁶ | X | Base |
| H | Н | Н | CH ₃ | S | 2-(N,N-diacetyl)- |
| | | : | | | guanine |
| Н | H | Н | CH ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| Н | H | H | CH ₃ | S | 8-fluoroguanine |
| Н | Н | H | CH ₃ | S | guanine |
| Н | H | Н | CH ₃ | S | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| Н | H | H | CH ₃ | S | 2-fluoroadenine |
| Н | Н | Н | CH ₃ | S | 8-fluoroadenine |
| Н | Н | H | CH ₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| Н | H | Н | CH ₃ | S | adenine |
| monophosphate | Н | Н | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| monophosphate | Н | Н | CH ₃ | 0 | 6-O-acetyl |
| | | | | | guanine |
| monophosphate | H | Н | CH ₃ | 0 | 8-fluoroguanine |
| monophosphate | Н | Н | CH ₃ | 0 | guanine |
| monophosphate | H | H | CH ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| monophosphate | H | Н | CH ₃ | 0 | 2-fluoroadenine |
| monophosphate | H | Н | CH ₃ | 0 | 8-fluoroadenine |
| monophosphate | H | Н | CH ₃ | 0 | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | Н | Н | CH ₃ | 0 | adenine |
| monophosphate | Н | Н | CH ₃ | S | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| monophosphate | Н | Н | CH ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| monophosphate | Н | Н | CH ₃ | S | 8-fluoroguanine |
| monophosphate | H | Н | CH ₃ | S | guanine |
| | <u></u> | | | | |

| WO 01/90121 R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | R ⁶ | X | PCT/US01/16671 Base |
|-------------------------------|----------------|----------------|-----------------|----|------------------------|
| | Н | H | CH ₃ | S | 6-(N,N-diacetyl)- |
| monophosphate | п | 111 | CII3 | 3 | adenine |
| | | | | 1_ | |
| monophosphate | Н | H | CH ₃ | S | 2-fluoroadenine |
| monophosphate | H | H | CH₃ | S | 8-fluoroadenine |
| monophosphate | Н | H | CH₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | Н | H | CH ₃ | S | adenine |
| diphosphate | Н | Н | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| diphosphate | H | Н | CH ₃ | 0 | 6-O-acetyl |
| | | | | | guanine |
| diphosphate | Н | H | CH ₃ | 0 | 8-fluoroguanine |
| diphosphate | Н | H | CH ₃ | 0 | guanine |
| diphosphate | Н | H | CH ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| diphosphate | Н | Н | CH ₃ | 0 | 2-fluoroadenine |
| diphosphate | H . | H | CH ₃ | 0 | 8-fluoroadenine |
| diphosphate | H | H | CH ₃ | 0 | 2,8-difluoro- |
| | | | | | adenine |
| diphosphate | H | H | CH ₃ | 0 | adenine |
| diphosphate | H | H | CH ₃ | S | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| diphosphate | Н | H | CH ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| diphosphate | H | H | CH ₃ | S | 8-fluoroguanine |
| diphosphate | H | Н | CH ₃ | S | guanine |
| diphosphate | H | H | CH ₃ | S | 6-(N,N-diacetyl)- |
| • | | | | | adenine |
| diphosphate | H | Н | CH ₃ | S | 2-fluoroadenine |
| diphosphate | Н | H | CH ₃ | S | 8-fluoroadenine |
| diphosphate | Н | H | CH ₃ | S | 2,8-difluoro- |
| | | | | | adenine |

WO 01/90121 PCT/US01/16671 $\overline{\mathbf{R^6}}$ R^2 R^3 $\overline{\mathbf{X}}$ $\overline{\mathbf{R}^1}$ Base H CH₃ S adenine H diphosphate H CH₃ 2-(N,N-diacetyl)-0 triphosphate H guanine CH₃ 0 6-O-acetyl H H triphosphate guanine H CH₃ 0 8-fluoroguanine H triphosphate H CH₃ triphosphate H 0 guanine CH₃ 6-(N,N-diacetyl)-Н 0 Η triphosphate adenine CH₃ O 2-fluoroadenine triphosphate Η Η 8-fluoroadenine CH₃ triphosphate Η $\overline{\mathbf{H}}$ O 2,8-difluorotriphosphate H H CH_3 adenine 2-(N,N-diacetyl)-Η CH_3 0 triphosphate H guanine S 6-O-acetyl H Η CH_3 triphosphate guanine S 8-fluoroguanine Н CH₃ Н triphosphate guanine H H CH_3 S triphosphate 6-(N,N-diacetyl)-H CH₃ S triphosphate Н adenine CH₃ 2-fluoroadenine triphosphate H H S triphosphate Η H CH_3 S 8-fluoroadenine CH₃ S 2,8-difluoro-Н H triphosphate adenine S H CH₃ adenine H triphosphate 2-(N,N-diacetyl)-O monophosphate monophosphate monophosphate CF_3 guanine 6-O-acetyl monophosphate monophosphate monophosphate CF_3 0 guanine monophosphate monophosphate monophosphate CF₃ 0 8-fluoroguanine monophosphate monophosphate CF₃ 0 guanine monophosphate

| WO 01/90121 | | | | | PCT/US01/16671 |
|----------------|----------------|----------------|-----------------|---|-------------------|
| R ¹ | R ² | R ³ | R ⁶ | X | Base |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,8-difluoro- |
| | | · | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 8-fluoroguanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 8-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | adenine |
| acetyl | acetyl | acetyl | CF ₃ | 0 | guanine |
| acetyl | acetyl | acetyl | CF ₃ | S | guanine |
| acetyl | acetyl | acetyl | 2-bromo- | 0 | guanine |
| | | | vinyl | | |
| acetyl | acetyl | acetyl | 2-bromo- | S | guanine |
| | | | vinyl | | |

Alternatively, the following nucleosides of Formula XI are prepared, using the appropriate sugar and pyrimidine or purine bases.

wherein:

| R ¹ | R ² | R ⁷ | \mathbb{R}^6 | X | Base |
|----------------|----------------|----------------|-----------------|---|--------------------------|
| H | Н | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| H | H | Н | CH ₃ | 0 | Hypoxanthine |
| Н | H | Н | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| H | H | Н | CH ₃ | 0 | Thymine |
| H | Н | Н | CH ₃ | 0 | Cytosine |
| H | Н | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| H | Н | Н | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| H | Н | Н | CH ₃ | 0 | Uracil |
| H | Н | Н | CH ₃ | 0 | 5-Fluorouracil |
| H | Н | Н | CH ₃ | S | 2,4-O-Diacetyluracil |
| H | Н | H | CH ₃ | S | Hypoxanthine |
| H | Н | Н | CH ₃ | S | 2,4-O-Diacetylthymine |
| H | Н | Н | CH ₃ | S | Thymine |
| H | H | Н | CH ₃ | S | Cytosine |
| H | Н | Н | CH ₃ | S | 4-(N-mono-acetyl)cytosin |
| H | Н | H | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| H | Н | Н | CH ₃ | S | Uracil |
| H | Н | Н | CH ₃ | S | 5-Fluorouracil |
| | | | CH ₃ | | |
| monophosphate | Н | H | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | Н | H | CH ₃ | 0 | Hypoxanthine |
| monophosphate | Н | H | CH ₃ | 0 | 2,4-O-Diacetylthymine |

WO 01/90121 R¹ R⁷ R⁶ X R² Base

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| | | | | | [|
|---------------|---|---|-----------------|---|--------------------------|
| monophosphate | Н | Н | CH ₃ | 0 | Thymine |
| monophosphate | Н | H | CH ₃ | 0 | Cytosine |
| monophosphate | Н | H | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| monophosphate | Н | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | H | Н | CH ₃ | 0 | Uracil |
| monophosphate | H | H | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | Н | H | CH ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | Н | H | CH ₃ | S | Hypoxanthine |
| monophosphate | Н | H | CH ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | Н | Н | CH ₃ | S | Thymine |
| monophosphate | H | H | CH ₃ | S | Cytosine |
| monophosphate | Н | Н | CH ₃ | S | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| monophosphate | Н | Н | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | Н | Н | CH ₃ | S | Uracil |
| monophosphate | Н | H | CH ₃ | S | 5-Fluorouracil |
| diphosphate | H | H | CH ₃ | 0 | 2,4-O-Diacetylurac |
| diphosphate | Н | H | CH ₃ | 0 | Hypoxanthine |
| diphosphate | Н | H | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| diphosphate | Н | H | CH ₃ | 0 | Thymine |
| diphosphate | Н | H | CH ₃ | 0 | Cytosine |
| diphosphate | Н | H | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| diphosphate | Н | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| diphosphate | Н | H | CH ₃ | 0 | Uracil |
| diphosphate | H | H | CH ₃ | 0 | 5-Fluorouracil |
| diphosphate | H | H | CH ₃ | S | 2,4-O-Diacetyluracil |
| diphosphate | Н | H | CH ₃ | S | Hypoxanthine |
| diphosphate | Н | H | CH ₃ | S | 2,4-O-Diacetylthym |
| diphosphate | Н | H | CH ₃ | S | Thymine |

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|----------------------------|---------------------------------------|---------------------------------------|--|---|--|--|
| \mathbb{R}^2 | R^7 | R ⁶ | X | Base | | |
| Н | H | CH ₃ | S | Cytosine | | |
| Н | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil | | |
| Н | H | CH ₃ | 0 | Hypoxanthine | | |
| Н | H | CH ₃ | 0 | 2,4-O-Diacetylthymine | | |
| Н | H | CH ₃ | 0 | Thymine | | |
| H | H | CH ₃ | 0 | Cytosine | | |
| H | H | CH ₃ | 0 | 4-(N-mono- | | |
| | | | | acetyl)cytosine | | |
| H | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytos | | |
| Н | H | CH ₃ | 0 | Uracil | | |
| H | Н | CH ₃ | 0 | 5-Fluorouracil | | |
| H | H | CH ₃ | S | 2,4-O-Diacetyluracil | | |
| Н | H | CH ₃ | S | Hypoxanthine | | |
| Н | Н | CH ₃ | S | 2,4-O-Diacetylthym | | |
| Н | Н | CH ₃ | S | Thymine | | |
| Н | H | CH ₃ | S | Cytosine | | |
| monophosphate | Br | CF ₃ | 0 | 2,4-O-Diacetyluracil | | |
| monophosphate | Br | CF ₃ | 0 | Hypoxanthine | | |
| monophosphate | Br | CF ₃ | 0 | 2,4-O-Diacetylthymine | | |
| monophosphate | Br | CF ₃ | 0 | Thymine | | |
| monophosphate | Br | CF ₃ | 0 | Cytosine | | |
| monophosphate | Br | CF ₃ | 0 | 4-(N-mono- | | |
| | | | | acetyl)cytosine | | |
| monophosphate | Br | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine | | |
| monophosphate | Br | CF ₃ | 0 | Uracil | | |
| monophosphate | Br | CF ₃ | 0 | 5-Fluorouracil | | |
| monophosphate | Br | CF ₃ | S | 2,4-O-Diacetyluracil | | |
| monophosphate | Br | CF ₃ | S | Hypoxanthine | | |
| monophosphate | Br | CF ₃ | S | 2,4-O-Diacetylthymine | | |
| monophosphate | Br | CF ₃ | S | Thymine | | |
| monophosphate | Br | CF ₃ | S | Cytosine | | |
| | H H H H H H H H H H H H H H H H H H H | H H H H H H H H H H H H H H H H H H H | H H H CH3 H CH3 H CH3 H H CH3 H CH | H H CH3 S H H CH3 O H H CH3 S M H CH3 S M H CH3 S M H CH3 S M H CH3 S <tr< td=""></tr<> | | |

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|------------------------|----------------|----------------|-------------------|---|-------------------------------|--|
| \mathbb{R}^1 | \mathbb{R}^2 | \mathbb{R}^7 | \mathbb{R}^6 | X | Base | |
| monophosphate | monophosphate | Br | CF ₃ | S | 4-(N-mono- acetyl)cytosine | |
| monophosphate | monophosphate | Br | CF ₃ | S | 4-(N,N-diacetyl)cytos | |
| monophosphate | monophosphate | Br | CF ₃ | S | Uracil | |
| monophosphate | monophosphate | Br | CF ₃ | S | 5-Fluorouracil | |
| acetyl | acetyl | NO2 | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine | |
| acetyl | acetyl | NO2 | CF ₃ | S | 4-(N,N-diacetyl)cytosine | |
| acetyl | acetyl | NO2 | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine | |
| acetyl | acetyl | NO2 | 2-bromo- vinyl | S | 4-(N,N-diacetyl)cytosine | |

Alternatively, the following nucleosides of Formula XII are prepared, using the appropriate sugar and pyrimidine or purine bases.

wherein:

| \mathbb{R}^1 | R ⁶ | X | Base | |
|----------------|-----------------|---|---------------------------|--|
| Н | CH ₃ | О | 2,4-O-Diacetyluracil | |
| H | CH ₃ | 0 | Hypoxanthine | |
| H | CH ₃ | 0 | 2,4-O-Diacetylthymine | |
| H | CH ₃ | 0 | Thymine | |
| Н | CH ₃ | 0 | Cytosine | |
| H | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine | |
| H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine | |
| H | CH ₃ | 0 | Uracil | |
| H | CH ₃ | 0 | 5-Fluorouracil | |

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|---------------|-----------------|-----|---------------------------|
| R. | R ⁶ | X | Base |
| H | CH ₃ | S | 2,4-O-Diacetyluracil |
| H | CH ₃ | S | Hypoxanthine |
| Н | CH ₃ | S | 2,4-O-Diacetylthymine |
| Н | CH ₃ | S | Thymine |
| Н | CH ₃ | S | Cytosine |
| Н | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| H | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| Н | CH ₃ | S | Uracil |
| Н | CH ₃ | S | 5-Fluorouracil |
| monophosphate | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | CH ₃ | 0 | Hypoxanthine |
| monophosphate | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| monophosphate | CH ₃ | 0 | Thymine |
| monophosphate | CH ₃ | 0 | Cytosine |
| monophosphate | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | CH ₃ | 0 | Uracil |
| monophosphate | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | CH ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | CH ₃ | S | Hypoxanthine |
| monophosphate | CH ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | CH ₃ | S | Thymine |
| monophosphate | CH ₃ | S | Cytosine |
| monophosphate | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | CH ₃ | S | Uracil |
| monophosphate | CH ₃ | S | 5-Fluorouracil |
| diphosphate | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| diphosphate | CH ₃ | 0 | Hypoxanthine |
| diphosphate | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| diphosphate | CH ₃ | 0 | Thymine |

| | | PCT |
|-----------------|---|---|
| R ⁶ | X | Base |
| CH ₃ | 0 | Cytosine |
| CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| CH ₃ | 0 | Uracil |
| CH ₃ | 0 | 5-Fluorouracil |
| CH ₃ | S | 2,4-O-Diacetyluracil |
| CH ₃ | S | Hypoxanthine |
| CH ₃ | S | 2,4-O-Diacetylthymine |
| CH ₃ | S | Thymine |
| CH ₃ | S | Cytosine |
| CH ₃ | 0 | 2,4-O-Diacetyluracil |
| CH ₃ | 0 | Hypoxanthine |
| CH ₃ | 0 | 2,4-O-Diacetylthymine |
| CH ₃ | 0 | Thymine |
| CH ₃ | 0 | Cytosine |
| CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| CH ₃ | 0 | Uracil |
| CH ₃ | 0 | 5-Fluorouracil |
| CH ₃ | S | 2,4-O-Diacetyluracil |
| CH ₃ | S | Hypoxanthine |
| CH ₃ | S | 2,4-O-Diacetylthymine |
| CH ₃ | S | Thymine |
| CH ₃ | S | Cytosine |
| CF ₃ | 0 | 2,4-O-Diacetyluracil |
| CF ₃ | 0 | Hypoxanthine |
| CF ₃ | 0 | 2,4-O-Diacetylthymine |
| CF ₃ | 0 | Thymine |
| CF ₃ | 0 | Cytosine |
| CF ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| | CH ₃ | CH3 O CH3 O CH3 O CH3 O CH3 O CH3 S CH3 S CH3 S CH3 O CH3 S CF3 O CF3 O CF3 O CF3 O |

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|----------------|----------------|---|------|---------------------|
| \mathbb{R}^1 | R ⁶ | X | Rase | |

| R ¹ | R ⁶ | X | Base |
|----------------|-----------------|---|---------------------------|
| monophosphate | CF ₃ | 0 | Uracil |
| monophosphate | CF ₃ | 0 | 5-Fluorouracil |
| monophosphate | CF ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | CF ₃ | S | Hypoxanthine |
| monophosphate | CF ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | CF ₃ | S | Thymine |
| monophosphate | CF ₃ | S | Cytosine |
| monophosphate | CF ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | CF ₃ | S | Uracil |
| monophosphate | CF ₃ | S | 5-Fluorouracil |
| acetyl | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| acetyl | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| acetyl | 2-bromo-vinyl | 0 | 4-(N,N-diacetyl)cytosine |
| acetyl | 2-bromo-vinyl | S | 4-(N,N-diacetyl)cytosine |

Alternatively, the following nucleosides of Formula XVII are prepared, using the appropriate sugar and pyrimidine or purine bases.

$$R^{10}$$
 X R^{6} Base R^{9} R^{7} $(XVII)$

wherein:

| R ¹ | R ⁶ | \mathbf{R}^7 | X | Base | R ⁹ | R ¹⁰ |
|----------------|-----------------|----------------|---|-----------------------|----------------|-----------------|
| H | CH ₃ | H | 0 | 2,4-O-Diacetyluracil | NHAc | Me |
| H | CH ₃ | Н | 0 | Hypoxanthine | NH2 | Me |
| H | CH ₃ | H | 0 | 2,4-O-Diacetylthymine | NHAc | Me |
| H | CH ₃ | H | 0 | Thymine | NH2 | Me |

| \mathbb{R}^6 | \mathbb{R}^7 | X | | R ⁹ | R^{10} |
|-----------------|---|---|---|--|--|
| | | | | NH2 | Me |
| | | _ | | | Me |
| | | | 1 | <u> </u> | |
| | | | | | Me |
| | | _ | | | Me |
| CH ₃ | H | S | | | Me |
| CH ₃ | Н | S | Cytosine | NH2 | Me |
| CH ₃ | Н | S | 4-(N-mono-acetyl)cytosine | NHAc | Me |
| CH ₃ | Н | S | 4-(N,N-diacetyl)cytosine | NHAc | Me |
| CH ₃ | H | S | Uracil | NH2 | Me |
| CH ₃ | Н | S | 5-Fluorouracil | NH2 | Me |
| CH ₃ | Н | 0 | 2,4-O-Diacetyluracil | NHAc | Me |
| CH ₃ | Н | 0 | Hypoxanthine | NH2 | Me |
| CH ₃ | H | 0 | 2,4-O-Diacetylthymine | NHAc | Me |
| CH ₃ | Н | 0 | Thymine | NH2 | Me |
| CH ₃ | Н | 0 | Cytosine | NH2 | Me |
| CH ₃ | H | 0 | 4-(N-mono-acetyl)cytosine | NHAC | Me |
| CH ₃ | H | 0 | 4-(N,N-diacetyl)cytosine | NHAc | Me |
| CH ₃ | H | 0 | Uracil | NH2 | Me |
| CH ₃ | H | 0 | 5-Fluorouracil | NH2 | Me |
| CH ₃ | H | S | 2,4-O-Diacetyluracil | NHAc | Me |
| CH ₃ | \overline{H} | S | Hypoxanthine | NH2 | Me |
| CH ₃ | H | S | 2,4-O-Diacetylthymine | NHAc | Me |
| CH ₃ | H | S | Thymine | NH2 | Me |
| CH ₃ | H | S | | NH2 | Me |
| | H | S | | NHAc | Me |
| | H | S | | NHAc | Me |
| 1 | 1 | 1 - | (2.75) | 1 | |
| | CH ₃ | CH3 H CH3 H | CH3 H O CH3 H O CH3 H O CH3 H O CH3 H S CH3 H O CH3 H S CH3 H S <td< td=""><td>R⁶ R⁷ X Base CH₃ H O Cytosine CH₃ H O 4-(N-mono-acetyl)cytosine CH₃ H O 4-(N,N-diacetyl)cytosine CH₃ H O 5-Fluorouracil CH₃ H S 2,4-O-Diacetyluracil CH₃ H S Hypoxanthine CH₃ H S Thymine CH₃ H S Cytosine CH₃ H S Cytosine CH₃ H S Cytosine CH₃ H S 4-(N-mono-acetyl)cytosine CH₃ H S 5-Fluorouracil CH₃ H O 2,4-O-Diacetyluracil CH₃ H O 2,4-O-Diacetylthymine CH₃ H O 4-(N-mono-acetyl)cytosine CH₃ H O 4-(N-mono-acetyl)cytosine CH₃ H O 5-Fluorouracil <!--</td--><td>CH3 H O Cytosine NH2 CH3 H O 4-(N-mono-acetyl)cytosine NHAc CH3 H O 4-(N,N-diacetyl)cytosine NHAc CH3 H O Uracil NH2 CH3 H O 5-Fluorouracil NH2 CH3 H S 2,4-O-Diacetyluracil NHAc CH3 H S 14-O-Diacetyluracil NHAc CH3 H S 14-O-Diacetyluracil NHAc CH3 H S Cytosine NHAc CH3 H S Cytosine NHAc CH3 H S Uracil NHAc CH3 H S Uracil NHAc CH3 H S Uracil NHAc CH3 H O 2,4-O-Diacetyluracil NHAc CH3 H O Thymine NHAC CH3 H O C</td></td></td<> | R ⁶ R ⁷ X Base CH ₃ H O Cytosine CH ₃ H O 4-(N-mono-acetyl)cytosine CH ₃ H O 4-(N,N-diacetyl)cytosine CH ₃ H O 5-Fluorouracil CH ₃ H S 2,4-O-Diacetyluracil CH ₃ H S Hypoxanthine CH ₃ H S Thymine CH ₃ H S Cytosine CH ₃ H S Cytosine CH ₃ H S Cytosine CH ₃ H S 4-(N-mono-acetyl)cytosine CH ₃ H S 5-Fluorouracil CH ₃ H O 2,4-O-Diacetyluracil CH ₃ H O 2,4-O-Diacetylthymine CH ₃ H O 4-(N-mono-acetyl)cytosine CH ₃ H O 4-(N-mono-acetyl)cytosine CH ₃ H O 5-Fluorouracil </td <td>CH3 H O Cytosine NH2 CH3 H O 4-(N-mono-acetyl)cytosine NHAc CH3 H O 4-(N,N-diacetyl)cytosine NHAc CH3 H O Uracil NH2 CH3 H O 5-Fluorouracil NH2 CH3 H S 2,4-O-Diacetyluracil NHAc CH3 H S 14-O-Diacetyluracil NHAc CH3 H S 14-O-Diacetyluracil NHAc CH3 H S Cytosine NHAc CH3 H S Cytosine NHAc CH3 H S Uracil NHAc CH3 H S Uracil NHAc CH3 H S Uracil NHAc CH3 H O 2,4-O-Diacetyluracil NHAc CH3 H O Thymine NHAC CH3 H O C</td> | CH3 H O Cytosine NH2 CH3 H O 4-(N-mono-acetyl)cytosine NHAc CH3 H O 4-(N,N-diacetyl)cytosine NHAc CH3 H O Uracil NH2 CH3 H O 5-Fluorouracil NH2 CH3 H S 2,4-O-Diacetyluracil NHAc CH3 H S 14-O-Diacetyluracil NHAc CH3 H S 14-O-Diacetyluracil NHAc CH3 H S Cytosine NHAc CH3 H S Cytosine NHAc CH3 H S Uracil NHAc CH3 H S Uracil NHAc CH3 H S Uracil NHAc CH3 H O 2,4-O-Diacetyluracil NHAc CH3 H O Thymine NHAC CH3 H O C |

| WO 01/90121 R ¹ | R ⁶ | \mathbb{R}^7 | X | Base | 7/US01/166 R ⁹ | R ¹⁰ |
|-------------------------------|-----------------|----------------|---|---------------------------|------------------------------|-----------------|
| monophosphate | CH ₃ | H | S | 5-Fluorouracil | NH2 | Me |
| diphosphate | CH ₃ | H | 0 | 2,4-O-Diacetyluracil | NHAc | Me |
| diphosphate | CH ₃ | Н | Ō | Hypoxanthine | NH2 | Me |
| diphosphate | CH ₃ | H | 0 | 2,4-O-Diacetylthymine | NH2 | Me |
| diphosphate | CH ₃ | Н | 0 | Thymine | NH2 | Me |
| diphosphate | CH ₃ | H | 0 | Cytosine | NH2 | Me |
| diphosphate | CH ₃ | H | 0 | 4-(N-mono-acetyl)cytosine | NHAc | Me |
| diphosphate | CH ₃ | H | 0 | 4-(N,N-diacetyl)cytos | NHAc | Me |
| diphosphate | CH ₃ | H | 0 | Uracil | NH2 | Me |
| diphosphate | CH ₃ | H | 0 | 5-Fluorouracil | NH2 | Me |
| diphosphate | CH ₃ | H | S | 2,4-O-Diacetyluracil | NH2 | Me |
| diphosphate | CH ₃ | H | S | Hypoxanthine | NH2 | Me |
| diphosphate | CH ₃ | H | S | 2,4-O-Diacetylthymine | NHAc | Me |
| diphosphate | CH ₃ | H | S | Thymine | NH2 | Me |
| diphosphate | CH ₃ | H | S | Cytosine | NH2 | Me |
| triphosphate | CH ₃ | H | 0 | 2,4-O-Diacetyluracil | NHAc | Me |
| triphosphate | CH ₃ | H | 0 | Hypoxanthine | NHAc | Me |
| triphosphate | CH ₃ | H | 0 | 2,4-O-Diacetylthymine | NHAc | Me |
| triphosphate | CH ₃ | H | 0 | Thymine | NH2 | Me |
| triphosphate | CH ₃ | Н | 0 | Cytosine | NH2 | Me |
| triphosphate | CH ₃ | H | 0 | 4-(N-mono-acetyl)cytosine | NHAc | Me |
| triphosphate | CH ₃ | H | 0 | 4-(N,N-diacetyl)cytosine | NH2 | Me |
| triphosphate | CH ₃ | H | 0 | Uracil | NH2 | Me |
| triphosphate | CH ₃ | H | 0 | 5-Fluorouracil | NH2 | Me |
| triphosphate | CH ₃ | H | S | 2,4-O-Diacetyluracil | NH2 | Me |
| triphosphate | CH ₃ | H | S | Hypoxanthine | NH2 | Me |
| triphosphate | CH ₃ | H | S | 2,4-O-Diacetylthymine | NH2 | Me |
| triphosphate | CH ₃ | H | S | Thymine | NH2 | Me |
| triphosphate | CH ₃ | Н | S | Cytosine | NH2 | Me |
| monophosphate | CF ₃ | H | 0 | 2,4-O-Diacetyluracil | NH2 | Me |
| monophosphate | CF ₃ | H | 0 | Hypoxanthine | NH2 | Me |

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|---------------------------|-----------------|----------------|---|---------------------------|----------------|-----------------|
| \mathbb{R}^1 | R ⁶ | \mathbb{R}^7 | X | Base | R ⁹ | R ¹⁰ |
| monophosphate | CF ₃ | H | 0 | 2,4-O-Diacetylthymine | NH2 | Me |
| monophosphate | CF ₃ | H | 0 | Thymine | NH2 | Me |
| monophosphate | CF ₃ | Н | 0 | Cytosine | NH2 | Me |
| monophosphate | CF ₃ | H | 0 | 4-(N-mono-acetyl)cytosine | NH2 | Me |
| monophosphate | CF ₃ | Н | 0 | 4-(N,N-diacetyl)cytosine | NH2 | Me |
| monophosphate | CF ₃ | H | 0 | Uracil | NH2 | Me |
| monophosphate | CF ₃ | Н | 0 | 5-Fluorouracil | NH2 | Me |
| monophosphate | CF ₃ | H | S | 2,4-O-Diacetyluracil | NH2 | Me |
| monophosphate | CF ₃ | H | S | Hypoxanthine | NH2 | Me |
| monophosphate | CF ₃ | H | S | 2,4-O-Diacetylthymine | NH2 | Me |
| monophosphate | CF ₃ | Н | S | Thymine | NH2 | Me |
| monophosphate | CF ₃ | Н | S | Cytosine | NH2 | Me |
| monophosphate | CF ₃ | Н | S | 4-(N-mono-acetyl)cytosine | NH2 | Me |
| monophosphate | CF ₃ | Н | S | 4-(N,N-diacetyl)cytosine | NH2 | Me |
| monophosphate | CF ₃ | Н | S | Uracil | NH2 | Me |
| monophosphate | CF ₃ | H | S | 5-Fluorouracil | NH2 | Me |
| acetyl | CH ₃ | Н | 0 | 4-(N,N-diacetyl)cytosine | Н | Br |
| acetyl | CH ₃ | Н | S | 4-(N,N-diacetyl)cytosine | Н | Br |
| acetyl | CH ₃ | OH | 0 | 4-(N,N-diacetyl)cytosine | H | Br |
| acetyl | CH ₃ | OH | S | 4-(N,N-diacetyl)cytosine | H | Br |

Example 3: Preparation of 3'-C-methylriboadenine

The title compound can be prepared according to a published procedure (R.F. Nutt, M.J. Dickinson, F.W. Holly, and E. Walton, "Branched-chain sugar nucleosides. III. 3'-C-methyladenine", *J.Org. Chem.* 1968, 33, 1789-1795) (Scheme 9).

Scheme 9

(a) RuO₂ / NaIO₄; (b) MeMgI / TiCl₄; (c) HCl / MeOH / H₂O; (d) BzCl / pyridine; (e) AcBr, HBr / AcOH; (f) chloromercuri-6-benzamidopurine; (g) NH₃ / MeOH.

In a similar manner, but using the appropriate sugar and pyrimidine or purine bases, the following nucleosides of Formula III are prepared.

$$X^1$$
 N
 N
 X^2
 CH_3
 OR^2
 OR^3

(III)

wherein:

| \mathbb{R}^1 | R ² | \mathbb{R}^3 | X ¹ | X ² | Y |
|----------------|----------------|----------------|----------------|----------------|-----------------|
| Н | H | Н | H | H | Н |
| Н | H | Н | H | H | NH ₂ |
| Н | Н | Н | H | H | NH-cyclopropyl |
| Н | Н | H | H | H | NH-methyl |
| Н | Н | Н | H | H | NH-ethyl |
| Н | Н | Н | H | H | NH-acetyl |
| H | Н | Н | H | H | ОН |

| R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | X ² | Y |
|----------------|----------------|----------------|----------------|----------------|-----------------|
| H | Н | H | H | H | OMe |
| Н | H | H | H | H | OEt |
| H | Н | Н | Н | H | O-cyclopropyl |
| H | Н | H | H | H | O-acetyl |
| H | Н | H | H | H | SH |
| Н | Н | H | H | H | SMe |
| H | Н | H | H | H | SEt |
| H | Н | H | H | H | S-cyclopropyl |
| H | Н | H | H | H | F |
| Н | H | H | H | H | Cl |
| Н | H | Н | H | H | Br |
| Н | H | Н | H | H | I |
| monophosphate | H | H | H | H | NH ₂ |
| monophosphate | H | H | H | H | NH-acetyl |
| monophosphate | H | H | Н | H | NH-cyclopropyl |
| monophosphate | H | Н | H | H | NH-methyl |
| monophosphate | Н | H | Н | Н | NH-ethyl |
| monophosphate | H | H | H | Н | OH |
| monophosphate | Н | H | H | Н | O-acetyl |
| monophosphate | Н | Н | H | Н | OMe |
| monophosphate | Н | Н | Н | H | OEt |
| monophosphate | Н | Н | H | H | O-cyclopropyl |
| monophosphate | H | Н | H | Н | SH |
| monophosphate | Н | H | H | H | SMe |
| monophosphate | H | Н | Н | H | SEt |
| monophosphate | Н | H | Н | H | S-cyclopropyl |
| monophosphate | H | H | H | H | F |
| monophosphate | H | Н | H | H | Cl |
| monophosphate | H | Н | H | H | Br |
| monophosphate | Н | Н | H | H | I |
| diphosphate | H | H | H | H | NH ₂ |

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| R ¹ | R ² | \mathbb{R}^3 | X ¹ | X^2 | Y |
|----------------|----------------|----------------|----------------|-------|-----------------|
| | | | ļ | L | |
| diphosphate | Н | Н | H | H | NH-acetyl |
| diphosphate | Н | Н | H | H | NH-cyclopropyl |
| diphosphate | Н | Н | H | H | NH-methyl |
| diphosphate | Н | H | H | H | NH-ethyl |
| diphosphate | Н | Н | H | H | OH |
| diphosphate | Н | Н | H | H | O-acetyl |
| diphosphate | H | H | H | H | OMe |
| diphosphate | Н | Н | H | H | OEt |
| diphosphate | Н | Н | H | H | O-cyclopropyl |
| diphosphate | Н | Н | H | H | SH |
| diphosphate | H | Н | H | H | SMe |
| diphosphate | Н | Н . | Н | H | SEt |
| diphosphate | Н | Н | Н | Н | S-cyclopropyl |
| diphosphate | Н | Н | H | H | F |
| diphosphate | Н | Н | H | Н | Cl |
| diphosphate | Н | Н | H | H | Br |
| diphosphate | Н | Н | Н | H | I |
| triphosphate | Н | H | H | Н | NH ₂ |
| triphosphate | Н | Н | H | Н | NH-acetyl |
| triphosphate | Н | Н | Н | Н | NH-cyclopropyl |
| triphosphate | Н | H | Н | H | NH-methyl |
| triphosphate | Н | Н | H | Н | NH-ethyl |
| triphosphate | H | Н | H | Н | OH |
| triphosphate | Н | Н | H | Н | OMe |
| triphosphate | H | Н | H | H | OEt |
| triphosphate | H | Н | Н | H | O-cyclopropyl |
| triphosphate | H | Н | H | Н | O-acetyl |
| triphosphate | H | Н | Н | Н | SH |
| triphosphate | H | Н | Н | H | SMe |
| triphosphate | Н | Н | H | Н | SEt |
| triphosphate | Н | Н | H | Н | S-cyclopropyl |
| | | | | | |

PCT/US01/16671 $\overline{X^1}$ $\overline{X^2}$ $\overline{\mathbf{R^2}}$ $\overline{\mathbf{R}^3}$ $\overline{\mathbf{R}^{\mathbf{I}}}$ H F triphosphate H H H H H Cl triphosphate Η H H Η Η H Br triphosphate H H H I triphosphate H H NH₂ monophosphate monophosphate monophosphate H H H NH-cyclopropyl monophosphate monophosphate monophosphate H Η OH monophosphate monophosphate monophosphate F H Η monophosphate monophosphate monophosphate H H Cl monophosphate monophosphate monophosphate NH₂ Η H diphosphate diphosphate diphosphate H NH-cyclopropyl diphosphate diphosphate Η diphosphate diphosphate Η H OH diphosphate diphosphate Н Η F diphosphate diphosphate diphosphate Η Cl diphosphate Η diphosphate diphosphate NH₂ Н H triphosphate triphosphate triphosphate Η Η NH-cyclopropyl triphosphate triphosphate triphosphate triphosphate Η H OH triphosphate triphosphate F triphosphate triphosphate H Η triphosphate H Η Cl triphosphate triphosphate triphosphate F Η NH_2 Н \mathbf{H} Η F H H NH-cyclopropyl H H Н OH F Н H Η F F H H H H F Η Cl H H H H H H H Cl NH_2 Cl H NH-cyclopropyl H H Η Cl H OH H H Η F Čl Н H Η H Cl Cl H H H H H BrH NH_2 Η H H $\overline{\mathbf{H}}$ NH-cyclopropyl H Η Br

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|----------------|----------------|----------------|-----------------|----------------|-----------------|
| \mathbb{R}^1 | R ² | R ³ | X¹ | X ² | Y |
| Н | H | Н | Br | H | OH |
| Н | Н | Н | Br | Н | F |
| H | H | Н | Br | H | Cl |
| H | Н | Н | NH ₂ | H | NH ₂ |
| Н | H | Н | NH ₂ | H | NH-cyclopropyl |
| Н | H | Н | NH ₂ | Н | OH |
| Н | H | Н | NH ₂ | Н | F |
| H | H | Н | NH ₂ | H | Cl |
| H | H | Н | SH | Н | NH ₂ |
| H | H | Н | SH | H · | NH-cyclopropyl |
| Н | H | Н | SH | H | OH |
| H | H | H | SH | H | F |
| H | H | H | SH | Н | Cl |
| acetyl | H | H | H | Н | NH ₂ |
| acetyl | Н | H | Н | H | NH-cyclopropyl |
| acetyl | Н | Н | H | H | ОН |
| acetyl | H | H | H | H | F |
| acetyl | Н | Н | Н | H | Cl |
| acetyl | Н | Н | F | H | NH ₂ |
| acetyl | H | Н | F | H | NH-cyclopropyl |
| acetyl | H | Н | F | H | OH |
| acetyl | Н | Н | F | H | F |
| acetyl | H | Н | F | H | Cl |
| Н | acetyl | acetyl | Н | H | NH ₂ |
| Н | acetyl | acetyl | Н | Н | NH-cyclopropyl |
| Н | acetyl | acetyl | H | H | OH |
| Н | acetyl | acetyl | H | H | F |
| Н | acetyl | acetyl | H | H | Cl |
| acetyl | acetyl | acetyl | H | H | NH ₂ |
| acetyl | acetyl | acetyl | Н | H | NH-cyclopropyl |
| acetyl | acetyl | acetyl | H | Н | OH |
| | | | | | |

| R R R A A N acetyl acetyl acetyl H H F acetyl acetyl H H Cl monophosphate acetyl acetyl H H NH-cyclopropyl monophosphate acetyl acetyl H H OH monophosphate acetyl acetyl H H Cl monophosphate acetyl acetyl H H Cl diphosphate acetyl acetyl H H NH-cyclopropyl diphosphate acetyl acetyl H H NH-cyclopropyl diphosphate acetyl acetyl H H Cl diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H Cl triphosphate acetyl acetyl H H NH-cyclopropyl triphosphate | R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | X¹ | X^2 | Y |
|--|----------------|----------------|----------------|------------|-----------------|-----------------|
| | | | | j | ļ | |
| monophosphate acetyl acetyl H H NH2 monophosphate acetyl acetyl H H NH-cyclopropyl monophosphate acetyl acetyl H H H F monophosphate acetyl acetyl H H H CI diphosphate acetyl acetyl H H H NH-cyclopropyl diphosphate acetyl acetyl H H NH-cyclopropyl diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H F C diphosphate acetyl acetyl H H CI C triphosphate acetyl acetyl H H NH-cyclopropyl triphosphate acetyl acetyl H H CI <tr< td=""><td>acetyl</td><td></td><td>acetyl</td><td>ļ<u> </u></td><td></td><td></td></tr<> | acetyl | | acetyl | ļ <u> </u> | | |
| monophosphate acetyl acetyl H H NH-cyclopropyl monophosphate acetyl acetyl H H OH monophosphate acetyl acetyl H H F monophosphate acetyl acetyl H H H Cl diphosphate acetyl acetyl H H NH-cyclopropyl diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H F Cl diphosphate acetyl acetyl H H F Cl diphosphate acetyl acetyl H H Cl Cl triphosphate acetyl acetyl H H NH-cyclopropyl H H F Cl triphosphate acetyl acetyl H | acetyl | acetyl | acetyl | H | H | |
| monophosphate acetyl acetyl H H F monophosphate acetyl acetyl H H F monophosphate acetyl acetyl H H H CI diphosphate acetyl acetyl H H H NH-cyclopropyl diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H H CI triphosphate acetyl acetyl H H NH-cyclopropyl triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acet | monophosphate | acetyl | acetyl | H | H | NH ₂ |
| monophosphate acetyl acetyl H H H CI monophosphate acetyl acetyl H H H CI diphosphate acetyl acetyl H H H NH2 diphosphate acetyl acetyl H H H NH2 diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H H CI H H H H H H NH2 H CI H H H H H H NH2 NH2 H H H H NH2 NH-cyclopropyl H H H H H H NH2 OH H H H H H NH2 OH H H H H NH2 OOH H H H H H NH2 OO-cyclopropyl H H H H H NH2 O-cacetyl H H H H NH2 O-cacetyl H H H H NH2 SH | monophosphate | acetyl | acetyl | H | H | NH-cyclopropyl |
| monophosphate acetyl acetyl H H Cl diphosphate acetyl acetyl H H NH2 diphosphate acetyl acetyl H H NH-cyclopropyl diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H NH2 triphosphate acetyl acetyl H H NH-cyclopropyl triphosphate acetyl acetyl H H F | monophosphate | acetyl | acetyl | H | Н | OH . |
| diphosphate acetyl acetyl H H NH2 diphosphate acetyl acetyl H H NH-cyclopropyl diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H F H H F diphosphate acetyl acetyl H H H CI T T T T T CI T <td>monophosphate</td> <td>acetyl</td> <td>acetyl</td> <td>H</td> <td>H</td> <td>F</td> | monophosphate | acetyl | acetyl | H | H | F |
| diphosphate acetyl acetyl H H OH diphosphate acetyl acetyl H H H OH diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H H CI triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acetyl ACETYL H H H CI H H H H H H H NH2 H H H H H H NH2 NH2 H H H H H H NH2 NH-cyclopropyl H H H H H H H NH2 NH-methyl H H H H H H NH2 NH-methyl H H H H H H NH2 NH-cetyl H H H H H H NH2 OH H H H H H NH2 OH H H H H H NH2 OF H H H H H NH2 OF H H H H H H NH2 OF H H H H H NH2 O-cyclopropyl H H H H H NH2 NH2 O-cyclopropyl H H H H H NH2 NH2 SH | monophosphate | acetyl | acetyl | Н | H | Cl |
| diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H H F diphosphate acetyl acetyl H H H CI triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acetyl Acetyl H H H CI H H H H H H NH2 H H H H H NH2 NH-cyclopropyl H H H H H H NH2 NH-cyclopropyl H H H H H H H NH2 NH-cyclopropyl H H H H H H NH2 NH-cyclopropyl H H H H H H NH2 NH-cyclopropyl H H H H H H NH2 NH-cetyl H H H H H H NH2 OH H H H H H NH2 OH H H H H H NH2 OGEt H H H H H NH2 O-cyclopropyl H H H H H H NH2 O-cyclopropyl H H H H H NH2 O-cyclopropyl H H H H H NH2 O-cyclopropyl H H H H H NH2 NH2 O-cyclopropyl H H H H H NH2 NH2 SH | diphosphate | acetyl | acetyl | H | Н | NH ₂ |
| diphosphate acetyl acetyl H H F diphosphate acetyl acetyl H H H Cl triphosphate acetyl acetyl H H NH-cyclopropyl triphosphate acetyl acetyl H H OH triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H H Cl H H H H H NH NH | diphosphate | acetyl | acetyl | H | H | NH-cyclopropyl |
| diphosphate acetyl acetyl H H NH2 triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H NH2 triphosphate acetyl acetyl H H H NH4-cyclopropyl triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H H CI H H H H H NH2 H H H H H H NH2 NH4- H H H H H NH4 NH4 NH4- H H H H H NH4 OH H H H H H NH4 OH H H H H H NH4 OH H H H H NH4 OH H H H H H NH4 OH H H H H NH4 OFE H H H H H H NH4 OFE | diphosphate | acetyl | acetyl | H | H | ОН |
| triphosphate acetyl acetyl H H NH2 triphosphate acetyl acetyl H H H NH-cyclopropyl triphosphate acetyl acetyl H H H OH triphosphate acetyl acetyl H H H F triphosphate acetyl acetyl H H H F triphosphate acetyl ACI H H H H H H H NH2 H H H H H H NH2 H H H H H NH4 H H H H NH4 NH4 NH4 NH5 NH6 NH7 NH7 NH7 NH7 NH8 | diphosphate | acetyl | acetyl | H | H | F |
| triphosphate acetyl acetyl H H NH-cyclopropyl triphosphate acetyl acetyl H H OH triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H Cl H H H H H Cl H H H H H H H H H H H H H H H H NH-cyclopropyl H | diphosphate | acetyl | acetyl | H | Н | Cl |
| triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H Cl H H H H H Cl H H H H H H H H H H NH2 NH-ethyl H H H H NH2 OH H H H H NH2 OH H H H H NH2 O-cyclopropyl H H H H NH2 O-cyclopropyl H H H H NH2 O-cyclopropyl H | triphosphate | acetyl | acetyl | Н | H | NH ₂ |
| triphosphate acetyl acetyl H H F triphosphate acetyl acetyl H H Cl H H H H H H H H H H H H NH-2 NH-2 H H H H NH-2 NH-eyclopropyl H H H NH-2 NH-ethyl H H H NH-2 NH-ethyl H H H NH-2 NH-ethyl H H H NH-2 NH-acetyl H H H NH-2 OH H H H NH-2 OH H H H NH-2 Oet H H H NH-2 O-cyclopropyl H H H H NH-2 O-cyclopropyl H H H H NH-2 O-acetyl | triphosphate | acetyl | acetyl | H | Н | NH-cyclopropyl |
| triphosphate acetyl Acetyl H H Cl H H H H NH2 H H H H H NH2 NH-Cyclopropyl H H H H NH-Cyclopropyl H H H H NH-cyclopropyl H H < | triphosphate | acetyl | acetyl | H | Н | OH |
| H H H H H H NH2 H H H NH2 NH2 H H H H H NH2 NH-cyclopropyl H H H H H NH2 NH-methyl H H H H NH2 NH-methyl H H H H NH2 NH-acetyl H H H H H NH2 OH H H H NH2 OH H H H H NH2 OOH H H H H H NH2 OOEt H H H H H NH2 O-cyclopropyl H H H H H H NH2 SH H NH2 SH H H H H NH2 SH | triphosphate | acetyl | acetyl | H | H | F |
| H H H H NH2 NH2 H H H H NH-cyclopropyl H H H H NH-cyclopropyl H H H NH2 NH-methyl H H H NH2 NH-ethyl H H H NH2 NH-ethyl H H H NH2 OH H H H NH2 OH H H H NH2 Oet H H H NH2 O-cyclopropyl H H H NH2 O-acetyl H H H H NH2 SH H H H H NH2 SMe | triphosphate | acetyl | acetyl | Н | Н | CI |
| H H H H NH-cyclopropyl H H H H NH-cyclopropyl H H H NH- NH2 NH-methyl H H H H NH- NH2 NH-acetyl H H H H NH2 OH H H H H NH2 OH H H H H NH2 OEt H H H H NH2 O-cyclopropyl H H H H NH2 O-acetyl H H H NH2 SH H H H NH2 SMe | H | Н | Н | H | NH ₂ | Н |
| H H H H NH-methyl H H H NH- methyl NH- ethyl NH- ethyl H H NH- methyl H H NH- methyl H NH- NH2 OMe H | H | H | Н | H | NH ₂ | NH ₂ |
| H H H H NH-ethyl H H H H NH-acetyl H H H H NH2 OH H H H H NH2 OMe H H H H NH2 OEt H H H H NH2 O-cyclopropyl H H H NH2 O-acetyl H H H NH2 SH H H H NH2 SMe | H | Н | H | H | NH ₂ | NH-cyclopropyl |
| H H H NH2 NH-acetyl H H H H NH2 OH H H H H NH2 OMe H H H H NH2 OEt H H H NH2 O-cyclopropyl H H H NH2 O-acetyl H H H NH2 SH H H H NH2 SMe | H | Н | H | H | NH ₂ | NH-methyl |
| H H H H NH2 OH H H H H NH2 OMe H H H H NH2 OEt H H H NH2 O-cyclopropyl H H H NH2 O-acetyl H H H NH2 SH H H H NH2 SMe | H | Н | Н | Н | NH ₂ | NH-ethyl |
| H H H H NH2 OMe H H H H NH2 OEt H H H H NH2 O-cyclopropyl H H H NH2 O-acetyl H H H NH2 SH H H H NH2 SMe | H | Н | Н | H | NH ₂ | NH-acetyl |
| H H H H NH2 OEt H H H H NH2 O-cyclopropyl H H H H NH2 O-acetyl H H H NH2 SH H H H H NH2 SMe | H | Н | H | Н | NH ₂ | ОН |
| H H H H NH2 O-cyclopropyl H H H H NH2 O-acetyl H H H H NH2 SH H H H H NH2 SMe | H | Н | Н | H | NH ₂ | OMe |
| H H H H NH2 O-acetyl H H H H NH2 SH H H H H NH2 SMe | H | H | Н | H | NH ₂ | OEt |
| H H H H NH2 SH H H H H NH2 SMe | Н | Н | Н | Н | NH ₂ | O-cyclopropyl |
| H H H NH ₂ SMe | Н | Н | H | H | NH ₂ | O-acetyl |
| | H | Н | H | H | NH ₂ | SH |
| H H H NH ₂ SEt | H | Н | Н | H | NH ₂ | SMe |
| | H | Н | Н | H | NH ₂ | SEt |

| | | 11 | | 1/USU1/166/1 |
|----------------|---------------------------------------|---------------------------------------|---|--|
| R ² | R ³ | | | Y |
| Н | Н | Н | NH ₂ | S-cyclopropyl |
| Н | Н | H | NH ₂ | F |
| Н | Н | Н | NH ₂ | Cl |
| H | Н | H | NH ₂ | Br |
| Н | Н | Н | NH ₂ | I |
| Н | H | H | NH ₂ | NH ₂ |
| Н | Н | Н | NH ₂ | NH-acetyl |
| H. | Н | H | NH ₂ | NH-cyclopropyl |
| Н | Н | H | NH ₂ | NH-methyl |
| Н | Н | Н | NH ₂ | NH-ethyl |
| Н | Н | Н | NH ₂ | ОН |
| H | Н | Н | NH ₂ | O-acetyl |
| Н | Н | Н | NH ₂ | OMe |
| H | Н | Н | NH ₂ | OEt |
| Н | Н | H | NH ₂ | O-cyclopropyl |
| H | Н | Н | NH ₂ | SH |
| H | Н | H | NH ₂ | SMe |
| Н | Н | Н | NH ₂ | SEt |
| H | Н | H | NH ₂ | S-cyclopropyl |
| Н | Н | H | NH ₂ | F |
| Н | H | H | NH ₂ | Cl |
| Н | H , | Н | NH ₂ | Br |
| Н | Н | H | NH ₂ | I |
| Н | Н | Н | NH ₂ | NH ₂ |
| H | Н | H | NH ₂ | NH-acetyl |
| Н | Н | H | NH ₂ | NH-cyclopropyl |
| Н | H | H | NH ₂ | NH-methyl |
| Н | H | H | NH ₂ | NH-ethyl |
| H | H | H | NH ₂ | OH |
| H | H | H | NH ₂ | O-acetyl |
| H | Н | H | NH ₂ | OMe |
| | H H H H H H H H H H H H H H H H H H H | H H H H H H H H H H H H H H H H H H H | H H H H H < | R² R³ X¹ X² H H H NH2 H H H NH2 < |

| WO 01/90121 | | · · · · · · · · · · · · · · · · · · · | | | T/US01/16671 |
|----------------|----------------|---------------------------------------|----------------|-----------------|-----------------|
| R ¹ | R ² | \mathbb{R}^3 | X ¹ | X ² | Y |
| diphosphate | Н | Н | H | NH ₂ | OEt |
| diphosphate | H | Н | H | NH ₂ | O-cyclopropyl |
| diphosphate | H | Н | Н | NH ₂ | SH |
| diphosphate | H | H | Н | NH ₂ | SMe |
| diphosphate | H | Н | H | NH ₂ | SEt |
| diphosphate | Н | Н | H | NH ₂ | S-cyclopropyl |
| diphosphate | H | H | H | NH ₂ | F |
| diphosphate | Н | Н | H | NH ₂ | Cl . |
| diphosphate | H | H | H | NH ₂ | Br |
| diphosphate | H | H | H | NH ₂ | I |
| triphosphate | H | H | H | NH ₂ | NH ₂ |
| triphosphate | Н | Н | H | NH ₂ | NH-acetyl |
| triphosphate | H | Н | H | NH ₂ | NH-cyclopropyl |
| triphosphate | H | Н | H | NH ₂ | NH-methyl |
| triphosphate | H | H | H | NH ₂ | NH-ethyl |
| triphosphate | H | H | H | NH ₂ | ОН |
| triphosphate | H | Н | H | NH ₂ | OMe |
| triphosphate | Н | H | H | NH ₂ | OEt |
| triphosphate | H | H | H | NH ₂ | O-cyclopropyl |
| triphosphate | H | Н | H | NH ₂ | O-acetyl |
| triphosphate | Н | Н | H | NH ₂ | SH |
| triphosphate | Н | Н | H | NH ₂ | SMe |
| triphosphate | Н | Н | H | NH ₂ | SEt |
| triphosphate | Н | Н | H | NH ₂ | S-cyclopropyl |
| triphosphate | Н | Н | H | NH ₂ | F |
| triphosphate | Н | Н | H | NH ₂ | C1 |
| triphosphate | H | Н | H | NH ₂ | Br |
| triphosphate | Н | Н | H | NH ₂ | I . |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | NH ₂ |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | NH ₂ | ОН |
| | • | | | · | • |

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|-------------|----------------|
| | |

| R ¹ | \mathbb{R}^2 | R ³ | X ¹ | X^2 | Y |
|----------------|----------------|----------------|-----------------|-----------------|-----------------|
| monophosphate | monophosphate | monophosphate | H | NH ₂ | F |
| monophosphate | monophosphate | monophosphate | Н | NH ₂ | Cl |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | NH ₂ |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | OH |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | F |
| diphosphate | diphosphate | diphosphate | H | NH ₂ | Cl |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | NH ₂ |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | ОН |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | F |
| triphosphate | triphosphate | triphosphate | H | NH ₂ | Cl |
| Н | H | H | F | NH ₂ | NH ₂ |
| H | Н | Н | F | NH ₂ | NH-cyclopropyl |
| H | Н | H | F | NH ₂ | ОН |
| H | Н | H | F | NH ₂ | F |
| Н | Н | Н | F | NH ₂ | Cl |
| Н | Н | Н | Cl | NH ₂ | NH ₂ |
| H | Н | H | Cl | NH ₂ | NH-cyclopropyl |
| H | H | H | Cl | NH ₂ | ОН |
| H | Н | H | Cl | NH ₂ | F |
| H | H | Н | C1 | NH ₂ | Cl |
| Н | Н | Н | Br | NH ₂ | NH ₂ |
| H | H | H | Br | NH ₂ | NH-cyclopropyl |
| Н | H | Н | Br | NH ₂ | ОН |
| Н | H | Н | Br | NH ₂ | F |
| H | H | Н | Br | NH ₂ | Cl |
| Н | Н | H | NH ₂ | NH ₂ | NH ₂ |
| H | Н | H | NH ₂ | NH ₂ | NH-cyclopropyl |
| Н | Н | H | NH ₂ | NH ₂ | OH |
| H | Н | Н | NH ₂ | NH ₂ | F |

| \mathbb{R}^1 | R ² | R ³ | X ¹ | X ² | CT/US01/16671 Y |
|----------------|----------------|----------------|-----------------|-----------------|--------------------|
| H | H | Н | NH ₂ | NH ₂ | Cl |
| H | Н . | Н | SH | NH ₂ | NH ₂ |
| H | Н | H | SH | NH ₂ | NH-cyclopropyl |
| H | Н | H | SH | NH ₂ | OH |
| H | H | H | SH | NH ₂ | F |
| Н | H | Н | SH | NH ₂ | Cl |
| acetyl | H | H | H | NH ₂ | NH ₂ |
| acetyl | H | Н | H | NH ₂ | NH-cyclopropyl |
| acetyl | Н | H | Н | NH ₂ | OH |
| acetyl | H | H | H | NH ₂ | F |
| acetyl | H | Н | H | NH ₂ | Cl |
| acetyl | H | H | F | NH ₂ | NH ₂ |
| acetyl | Н | H | F | NH ₂ | NH-cyclopropyl |
| acetyl | Н | Н | F | NH ₂ | OH |
| acetyl | H | Н | F | NH ₂ | F |
| acetyl | H | Н | F | NH ₂ | Cl |
| H | acetyl | acetyl | H | NH ₂ | NH ₂ |
| H | acetyl | acetyl | Н | NH ₂ | NH-cyclopropyl |
| H | acetyl | acetyl | Н | NH ₂ | OH |
| H | acetyl | acetyl | H | NH ₂ | F |
| Н . | acetyl | acetyl | Н | NH ₂ | Cl |
| acetyl | acetyl | acetyl | H | NH ₂ | NH ₂ |
| acetyl | acetyl | acetyl | Н | NH ₂ | NH-cyclopropyl |
| acetyl | acetyl | acetyl | Н | NH ₂ | OH |
| acetyl | acetyl | acetyl | Н | NH ₂ | F |
| acetyl | acetyl | acetyl | H | NH ₂ | Cl |
| monophosphate | acetyl | acetyl | H | NH ₂ | NH ₂ |
| monophosphate | acetyl | acetyl | Н | NH ₂ | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | Н | NH ₂ | ОН |
| monophosphate | acetyl | acetyl | Н | NH ₂ | F |
| monophosphate | acetyl | acetyl | H | NH ₂ | Cl |

WO 01/90121 PCT/US01/16671 $\overline{X^1}$ \mathbb{R}^3 $\overline{X^2}$ $\overline{\mathbf{R^2}}$ $\overline{\mathbf{R}^{\mathrm{I}}}$ H NH₂ NH₂ diphosphate acetyl acetyl NH-cyclopropyl H NH_2 diphosphate acetyl acetyl NH₂ OH $\overline{\mathbf{H}}$ diphosphate acetyl acetyl F NH₂ acetyl H diphosphate acetyl Cl H NH_2 acetyl diphosphate acetyl NH₂ NH₂ H acetyl triphosphate acetyl NH-cyclopropyl H NH_2 acetyl triphosphate acetyl NH_2 OH H triphosphate acetyl acetyl F NH₂ H triphosphate acetyl acetyl Cl acetyl H NH_2 triphosphate acetyl H Cl H H H Η Cl Н Η H H Η Cl H $\overline{NH_2}$ H H H Cl NH-cyclopropyl H H H H Cl NH-methyl H Η Η H Cl NH-ethyl H H H H Cl NH-acetyl Η H H Η Н Cl OH H H Η Cl OMe H H Η H Cl OEt H Η H Η CI Η O-cyclopropyl H Н H C1 H H O-acetyl H H Cl SH H H H $\bar{\mathbf{H}}$ Cl H $\overline{\mathrm{H}}$ H SMe H Cl SEt H $\bar{\mathbf{H}}$ H H H H Cl S-cyclopropyl $\overline{\mathrm{H}}$ H Cl H NH₂ \overline{H} monophosphate Η

Η

H

H

 $\overline{\mathbf{H}}$

H

H

H

H

monophosphate

monophosphate

monophosphate

monophosphate

H

Η

H

Η

Cl

CI

Cl

Cl

NH-acetyl

NH-methyl

NH-ethyl

NH-cyclopropyl

| WO 01/90121 R ¹ | R ² | R ³ | X ¹ | X^2 | T/US01/16671 Y |
|-------------------------------|----------------|----------------|----------------|-------|-------------------|
| monophosphate | Н | H | H | Cl | OH |
| monophosphate | Н | H | H | Cl | O-acetyl |
| monophosphate | Н | Н | H | Cl | OMe |
| monophosphate | Н | H | H | Cl | OEt |
| monophosphate | Н | Н | H | Cl | O-cyclopropyl |
| monophosphate | Н | H | H | Cl | SH |
| monophosphate | Н | H | H | Cl | SMe |
| monophosphate | Н | H | Н | Cl | SEt |
| monophosphate | H | Н | H | Cl | S-cyclopropyl |
| diphosphate | Н | H | H | Cl | NH ₂ |
| diphosphate | H | H | H | Cl | NH-acetyl |
| diphosphate | Н | Н | Н | Cl | NH-cyclopropyl |
| diphosphate | Н | Н | Н | Cl | NH-methyl |
| diphosphate | Н | Н | Н | Cl | NH-ethyl |
| diphosphate | H | Н | Н | Cl | OH |
| diphosphate . | H | Н | Н | Cl | O-acetyl |
| diphosphate | H | H | Н | Cl | OMe |
| diphosphate | Н | Н | Н | Cl | OEt |
| diphosphate | Н | Н | Н | Cl | O-cyclopropyl |
| diphosphate | Н | Н | H | Cl | SH |
| diphosphate | H | Н | Н | Cl | SMe |
| diphosphate | H | Н | Н | Cl | SEt |
| diphosphate | Н | Н | H | Cl | S-cyclopropyl |
| triphosphate | Н | Н | Н | Cl | NH ₂ |
| triphosphate | Н | H | Н | Cl | NH-acetyl |
| triphosphate | Н | Н | H | Cl | NH-cyclopropyl |
| triphosphate | Н | Н | Н | Cl | NH-methyl |
| triphosphate | Н | Н | Н | Cl | NH-ethyl |
| triphosphate | H | Н | H | Cl | OH |
| triphosphate | Н | Н | H | Cl | OMe |
| triphosphate | H | Н | H | Cl | OEt |
| | | | | | |

| triphosphate H monophosphate monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate triphosphate | | WO 01/90121 PCT/US01/16671 | | | |
|--|----------------|----------------------------|----------------|-----------------|--|
| triphosphate H monophosphate monophosphate monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate diphosphate triphosphate t | R ³ | X ¹ | X ² | Y | |
| triphosphate H triphosphate H triphosphate H triphosphate H triphosphate H triphosphate H monophosphate monophosphate monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate diphosphate triphosphate triph | H | H | Cl | O-cyclopropyl | |
| triphosphate H triphosphate H triphosphate H monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate diphosphate triphosphate triphospha | H | Н | Cl | O-acetyl | |
| triphosphate H H H H H H H H H H H H H H H H H H H | Н | Н | Cl | SH | |
| triphosphate H monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate diphosphate triphosphate | Н | H | Cl | SMe | |
| monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate diphosphate diphosphate triphosphate tr | Н | H | Cl | SEt | |
| monophosphate monophosphate monophosphate diphosphate diphosphate diphosphate diphosphate diphosphate diphosphate triphosphate triphosp | Н | H | Cl | S-cyclopropyl | |
| monophosphate diphosphate diphosphate diphosphate diphosphate diphosphate diphosphate diphosphate diphosphate triphosphate | monophosphate | H | C1 | NH ₂ | |
| diphosphate diphosphate diphosphate diphosphate diphosphate diphosphate triphosphate triphosphat | monophosphate | Н | Cl | NH-cyclopropyl | |
| diphosphate diphosphate diphosphate diphosphate triphosphate triphosph | monophosphate | Н | Cl | OH | |
| diphosphatediphosphatedtriphosphatetriphosphatettriphosphatetriphosphatettriphosphatetriphosphatetHHH | diphosphate | H | Cl | NH ₂ | |
| triphosphate triph | diphosphate | H | Cl | NH-cyclopropyl | |
| triphosphate triph | diphosphate | H | Cl | OH | |
| triphosphate triphosphate the H H H H H H H H H H H H H H H H H H H | triphosphate | Н | Cl | NH ₂ | |
| H H H H H H H H H H H H H H H H H H H | triphosphate | H | Cl | NH-cyclopropyl | |
| H H H H H H H H H H H H H H H H H H H | triphosphate | Н | Cl | OH | |
| H H H H H H H H H H H H H H H H H H H | H | F | Cl | NH ₂ | |
| H H H H H H H H H H H H H H H H H H H | H | F | C1 | NH-cyclopropyl | |
| H H H H H H H H H H H H H H H H H H H | H | F | Cl | ОН | |
| H H H H H H H H H H H H H H H H H H H | H | Cl | Cl | NH ₂ | |
| H H H H H H H H H H H H H H H H H H H | Н | Cl | Cl | NH-cyclopropyl | |
| H H H H H H H H H H H H H H H H H H H | H | Cl | Cl | OH | |
| H H H F H H H H H H H H H H H H H H H H | H | Br | Cl | NH ₂ | |
| H H H H H H H H | H . | Br | Cl | NH-cyclopropyl | |
| H H H F H H | H | Br | Cl | OH | |
| H H H | Н | NH ₂ | Cl | NH ₂ | |
| H H F | H | NH ₂ | Cl | NH-cyclopropyl | |
| | H | NH ₂ | Cl | ОН | |
| H H F | Н | SH | C1 | NH ₂ | |
| 1 1 | Н | SH | Cl | NH-cyclopropyl | |
| H H H | H . | SH | Cl | OH | |
| acetyl H F | Н | H | Cl | NH ₂ | |

| WO 01/90121 PCT/US01/16671 | | | | | T/US01/16671 |
|----------------------------|----------------|----------------|----------------|-------|-----------------|
| R ¹ | R ² | \mathbb{R}^3 | X ¹ | X^2 | Y |
| acetyl | Н | H | H | Cl | NH-cyclopropyl |
| acetyl | H | Н | H | Cl | ОН |
| acetyl | H | Н | F | Cl | NH ₂ |
| acetyl | H | Н | F | Cl | NH-cyclopropyl |
| acetyl | Н | Н | F | Cl | OH |
| H | acetyl | acetyl | H | C1 | NH ₂ |
| H | acetyl | acetyl | H | C1 | NH-cyclopropyl |
| H | acetyl | acetyl | H | Cl | OH |
| acetyl | acetyl | acetyl | Н | Cl | NH ₂ |
| acetyl | acetyl | acetyl | Н | Cl | NH-cyclopropyl |
| acetyl | acetyl | acetyl | Н | Cl | ОН |
| monophosphate | acetyl | acetyl | H | C1 | NH ₂ |
| monophosphate | acetyl | acetyl | Н | C1 | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | H | Cl | OH |
| diphosphate | acetyl | acetyl | H | Cl | NH ₂ |
| diphosphate | acetyl | acetyl | Н | Cl | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | H | Cl | OH |
| triphosphate | acetyl | acetyl | Н | Cl | NH ₂ |
| triphosphate | acetyl | acetyl | Н | C1 | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | H | Cl | OH |
| Н | Н | Н | H | Cl | NH ₂ |
| H | H | Н | Н | Cl | NH-cyclopropyl |
| H | Н | Н | Н | Cl | ОН |
| Н | H | Н | Н | Br | NH ₂ |
| H | Н | Н | H | Br | NH-cyclopropyl |
| H | Н | Н | Н | Br | OH |

Alternatively, the following nucleosides of Formula VI are prepared, using the appropriate sugar and pyrimidine or purine bases.

$$X^{1}$$
 X^{1}
 X^{1

wherein:

| R ¹ | \mathbb{R}^2 | R ³ | X¹ | Y |
|----------------|----------------|----------------|----|-----------------|
| H | Н | Н | Н | Н |
| H | Н | Н | H | NH ₂ |
| Н | Н | Н | Н | NH-cyclopropyl |
| H | Н | Н | H | NH-methyl |
| Н | Н | Н | H | NH-ethyl |
| H | Н | Н | H | NH-acetyl |
| H | H | H | H | OH |
| H | Н | H | H | OMe |
| H | Н | Н | Н | OEt |
| H | Н | Н | H | O-cyclopropyl |
| H | Н | Н | H | O-acetyl |
| H | Н . | Н | H | SH |
| H | Н | Н | H | SMe |
| H | Н | Н | H | SEt |
| Н | Н | Н | H | S-cyclopropyl |
| monophosphate | Н | H | H | NH ₂ |
| monophosphate | H | Н | H | NH-acetyl |
| monophosphate | Н | Н | H | NH-cyclopropyl |
| monophosphate | H | Н | H | NH-methyl |
| monophosphate | Н | H | H | NH-ethyl |
| monophosphate | H | Н | Н | OH |
| monophosphate | H | Н | H | O-acetyl |

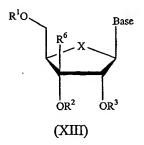
| WO 01/90121 | | | | PCT/US01/166' |
|----------------|----------------|----------------|----------------|-----------------|
| \mathbb{R}^1 | \mathbb{R}^2 | \mathbb{R}^3 | X ¹ | Y |
| monophosphate | H | Н | Н | OMe |
| monophosphate | H | H | H | OEt |
| monophosphate | Н | H | H | O-cyclopropyl |
| monophosphate | Н | Н | Н | SH |
| monophosphate | H | Н | Н | SMe |
| monophosphate | H | Н | H | SEt |
| monophosphate | Н | Н | Н | S-cyclopropyl |
| diphosphate | Н | Н | H | NH ₂ |
| diphosphate | Н | Н | H | NH-acetyl |
| diphosphate | Н | H | H | NH-cyclopropyl |
| diphosphate | H | Н | Н | NH-methyl |
| diphosphate | H | Н | H | NH-ethyl |
| diphosphate | Н | Н | H | ОН |
| diphosphate | H | Н | Н | O-acetyl |
| diphosphate | H | Н | Н | OMe |
| diphosphate | H | H | Н | OEt |
| diphosphate | Н | Н | Н | O-cyclopropyl |
| diphosphate | H | Н | Н | SH |
| diphosphate | Н | Н | Н | SMe |
| diphosphate | H | H | H | SEt |
| diphosphate | H | H | H | S-cyclopropyl |
| triphosphate | H | H | Н | NH ₂ |
| triphosphate | Н | H | H | NH-acetyl |
| triphosphate | H | H | H | NH-cyclopropyl |
| triphosphate | H | H | H | NH-methyl |
| triphosphate | H | H | Н | NH-ethyl |
| triphosphate | Н | H | H | ОН |
| triphosphate | H | H | Н | OMe |
| triphosphate | H | Н | Н | OEt |
| triphosphate | H | H | H | O-cyclopropyl |
| triphosphate | Н | Н | Н | O-acetyl |

| WO 01/90121 | | | | PCT/US01/1667 |
|----------------|----------------|----------------|-----------------|-----------------|
| R ¹ | R ² | \mathbb{R}^3 | X¹ | Y |
| triphosphate | Н | H | H | SH |
| triphosphate | H | H | Н | SMe |
| triphosphate | H | Н | Н | SEt |
| triphosphate | Н | Н | Н | S-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | NH ₂ |
| monophosphate | monophosphate | monophosphate | H | NH-cyclopropyl |
| monophosphate | monophosphate | monophosphate | H | ОН |
| diphosphate | diphosphate | diphosphate | H | NH ₂ |
| diphosphate | diphosphate | diphosphate | Н | NH-cyclopropyl |
| diphosphate | diphosphate | diphosphate | H | ОН |
| triphosphate | triphosphate | triphosphate | H | NH ₂ |
| triphosphate | triphosphate | triphosphate | Н | NH-cyclopropyl |
| triphosphate | triphosphate | triphosphate | H | OH |
| H | Н | H | F | NH ₂ |
| Н | Н | H | F | NH-cyclopropyl |
| Н | Н | Н | F | ОН |
| Н | Н | Н | Cl | NH ₂ |
| Н | Н | H | C1 | NH-cyclopropyl |
| Н | Н | H | Cl | OH |
| H | H | Н | Br | NH ₂ |
| Н | Н | H | Br | NH-cyclopropyl |
| Н | Н | Н | Br | ОН |
| H | Н | Н | NH ₂ | NH ₂ |
| H | Н | H | NH ₂ | NH-cyclopropyl |
| H | Н | Н | NH ₂ | ОН |
| Н | H | H | SH | NH ₂ |
| H | Н | Н | SH | NH-cyclopropyl |
| H | Н | H | SH | ОН |
| acetyl | Н | Н | H | NH ₂ |
| acetyl | Н | Н | H | NH-cyclopropyl |
| acetyl | Н | H | H | ОН |

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|-------------|------|-------|-----------|
| | | | |

| \mathbb{R}^{1} | R ² | R ³ | X ¹ | Y |
|------------------|----------------|----------------|----------------|-----------------|
| acetyl | H | H | F | NH ₂ |
| acetyl | H | Ĥ | F | NH-cyclopropyl |
| acetyl | Н | H | F | ОН |
| H | acetyl | acetyl | Н | NH ₂ |
| H | acetyl | acetyl | Н | NH-cyclopropyl |
| H | acetyl | acetyl | Н | ОН |
| acetyl | acetyl | acetyl | Н | NH ₂ |
| acetyl | acetyl | acetyl | Н | NH-cyclopropyl |
| acetyl | acetyl | acetyl | H | ОН |
| monophosphate | acetyl | acetyl | Н | NH ₂ |
| monophosphate | acetyl | acetyl | H | NH-cyclopropyl |
| monophosphate | acetyl | acetyl | Н | ОН |
| diphosphate | acetyl | acetyl | Н | NH ₂ |
| diphosphate | acetyl | acetyl | H | NH-cyclopropyl |
| diphosphate | acetyl | acetyl | H | ОН |
| triphosphate | acetyl | acetyl | H | NH ₂ |
| triphosphate | acetyl | acetyl | Н | NH-cyclopropyl |
| triphosphate | acetyl | acetyl | Н | ОН |

Alternatively, the following nucleosides of Formula XIII are prepared, using the appropriate sugar and pyrimidine or purine bases.



wherein:

| \mathbb{R}^1 | R ² | R ³ | R ⁶ | X | Base |
|----------------|----------------|----------------|-----------------|---|----------------|
| Н | H | Н | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetyluracil |
| H | H | H | CH ₃ | О | Hypoxanthine |

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|----------------------------|----------------|----------------|-----------------|---|-------------------|
| R ¹ | R ² | \mathbb{R}^3 | R ⁶ | X | Base |
| H | Н | Н | CH ₃ | 0 | 2,4-0- |
| | | | | | Diacetylthymine |
| H | Н | H | CH ₃ | 0 | Thymine |
| H | H | Н | CH ₃ | 0 | Cytosine |
| H | Н | Н | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |
| H | Н | Н | CH ₃ | 0 | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| H | Н | Н | CH ₃ | 0 | Uracil |
| H | Н | Н | CH ₃ | 0 | 5-Fluorouracil |
| H | H | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetyluraci |
| H | Н | Н | CH ₃ | S | Hypoxanthine |
| H | Н | Н | CH ₃ | S | 2,4-O- |
| | | | | | Diacetylthymine |
| H | Н | Н | CH ₃ | S | Thymine |
| H | Н | Н | CH ₃ | S | Cytosine |
| H | Н | Н | CH ₃ | S | 4-(N-mono- |
| : | | | | | acetyl)cytosine |
| H | Н | Н | CH ₃ | S | 4-(N,N- |
| | | | | | diacetyl)cytosine |
| H | H | ·H | CH ₃ | S | Uracil |
| H | H | Н | CH ₃ | S | 5-Fluorouracil |
| monophosphate | H | H | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetyluracil |
| monophosphate | Н | Н | CH ₃ | 0 | Hypoxanthine |
| monophosphate | Н | Н | CH ₃ | 0 | 2,4-O- |
| | | | | | Diacetylthym |
| monophosphate | H | Н | CH ₃ | 0 | Thymine |
| monophosphate | Н | Н | CH ₃ | 0 | Cytosine |
| monophosphate | Н | H | CH ₃ | 0 | 4-(N-mono- |
| | | | | | acetyl)cytosine |

R⁶ $\overline{\mathbf{R^2}}$ $\overline{\mathbf{R}^1}$ R³ X Base H CH₃ 4-(N,N-H ō monophosphate diacetyl)cytosine H CH₃ Uracil H Ō monophosphate CH₃ H H 0 5-Fluorouracil monophosphate H H CH₃ S 2,4-0monophosphate Diacetyluracil H H CH₃ S Hypoxanthine monophosphate 2,4-0-H H CH₃ S monophosphate Diacetylthym CH₃ Н H S Thymine monophosphate CH₃ Cytosine H monophosphate $\overline{\mathbf{H}}$ S H H CH₃ 4-(N-monomonophosphate acetyl)cytosine S 4-(N,N-H H CH₃ monophosphate diacetyl)cytosine CH₃ Uracil monophosphate H H S H CH₃ 5-Fluorouracil H S monophosphate 2,4-0-Η Η CH_3 O diphosphate Diacetyluracil H H CH₃ O Hypoxanthine diphosphate diphosphate H Η CH_3 O 2,4-0-Diacetylthymine diphosphate Η H CH_3 O Thymine CH₃ Cytosine H H 0 diphosphate 4-(N-monodiphosphate H Η CH_3 0 acetyl)cytosine 4-(N,N- $\overline{\mathbf{H}}$ CH_3 O diphosphate H diacetyl)cytosine Uracil H H CH_3 0 diphosphate CH₃ 5-Fluorouracil H diphosphate H 0 diphosphate H Η CH₃ S 2,4-0-Diacetyluracil

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WO 01/90121 PCT/US01/16671 $\overline{\mathbf{R}^1}$ $\overline{\mathbf{R}^2}$ R^3 R⁶ $\overline{\mathbf{X}}$ Base diphosphate H Η CH₃ Hypoxanthine S Η S 2,4-0diphosphate Η CH_3 Diacetylthym Η CH₃ S Thymine diphosphate Η H CH₃ S Cytosine diphosphate H Η H CH₃ O 2,4-0triphosphate Diacetyluracil Hypoxanthine H H CH_3 O triphosphate CH₃ 2,4-0-H H 0 triphosphate Diacetylthymine H CH₃ O. Thymine Η triphosphate CH₃ 0 Cytosine triphosphate H Η H Η CH_3 O 4-(N-monotriphosphate acetyl)cytosine H Н CH₃ O 4-(N,Ntriphosphate diacetyl)cytosine CH₃ Uracil Η H O triphosphate 5-Fluorouracil H CH_3 O H triphosphate CH_3 2,4-0triphosphate Η Η S Diacetyluracil Hypoxanthine CH_3 S triphosphate H Η CH₃ S 2,4-0-H triphosphate $\overline{\mathbf{H}}$ Diacetylthymine CH₃ Thymine H Η S triphosphate Η CH₃ S Cytosine triphosphate Η 2,4-0-CF₃ O monophosphate monophosphate monophosphate Diacetyluracil CF₃ O Hypoxanthine monophosphate monophosphate monophosphate 2,4-0-CF₃ monophosphate O monophosphate monophosphate Diacetylthymine O CF₃ Thymine monophosphate monophosphate monophosphate CF₃ O Cytosine monophosphate monophosphate monophosphate

 $\overline{\mathbf{R^6}}$ R^1 $\overline{\mathbb{R}^2}$ R^3 X Base 4-(N-monomonophosphate monophosphate monophosphate CF₃ 0 acetyl)cytosine CF₃ 0 4-(N,Nmonophosphate monophosphate monophosphate diacetyl)cytosine monophosphate monophosphate CF₃ 0 Uracil monophosphate CF₃ 0 5-Fluorouracil monophosphate monophosphate monophosphate monophosphate CF₃ S 2,4-0monophosphate monophosphate Diacetyluracil CF₃ monophosphate S Hypoxanthine monophosphate monophosphate S 2,4-0- CF_3 monophosphate monophosphate monophosphate Diacetylthymine monophosphate CF₃ S Thymine monophosphate monophosphate CF₃ Cytosine monophosphate monophosphate monophosphate S S monophosphate monophosphate CF₃ 4-(N-monomonophosphate acetyl)cytosine CF₃ S 4-(N,Nmonophosphate monophosphate monophosphate diacetyl)cytosine CF₃ S Uracil monophosphate monophosphate monophosphate 5-Fluorouracil CF_3 S monophosphate monophosphate monophosphate CF₃ O 4-(N,Nacetyl acetyl acetyl diacetyl)cytosine S 4-(N,N-CF₃ acetyl acetyl acetyl diacetyl)cytosine 4-(N,Nacetyl 2-bromo-0 acetyl acetyl vinyl diacetyl)cytosine acetyl 2-bromo-S 4-(N,Nacetyl acetyl diacetyl)cytosine vinyl CH₃ 2-(N,N-diacetyl)-Н Η H guanine $\overline{\mathbf{H}}$ CH_3 0 6-O-acetyl H H guanine CH₃ O 8-fluoroguanine Η Η H

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WO 01/90121 PCT/US01/16671 $\overline{\mathbf{R}^3}$ R⁶ $\overline{R^1}$ R^2 $\overline{\mathbf{X}}$ Base CH₃ $\overline{\mathbf{H}}$ H H 0 guanine 6-(N,N-diacetyl)-H H H CH₃ Ō adenine CH₃ H \overline{o} 2-fluoroadenine H Η CH₃ H H H 0 8-fluoroadenine CH₃ H 2,8-difluoro-H H Ō adenine CH₃ H o adenine H H 2-(N,N-diacetyl)-CH₃ H H Η S guanine 6-O-acetyl H H CH_3 S Η guanine 8-fluoroguanine CH₃ S H H H CH₃ guanine H S \overline{H} H 6-(N,N-diacetyl)-H H Η CH₃ S adenine $\overline{\mathbf{H}}$ CH₃ S 2-fluoroadenine H Н H CH₃ S 8-fluoroadenine H H CH₃ 2,8-difluoro-H S Η H adenine adenine H CH₃ S H H CH₃ H 2-(N,N-diacetyl)-0 monophosphate H guanine 6-O-acetyl $\overline{\mathbf{H}}$ H CH₃ O monophosphate guanine H H CH₃ $\overline{0}$ 8-fluoroguanine monophosphate CH₃ H O guanine H monophosphate CH₃ 6-(N,N-diacetyl)-H H O monophosphate adenine CH₃ ō 2-fluoroadenine H H monophosphate CH₃ 0 8-fluoroadenine monophosphate H H

WO 01/90121 PCT/US01/16671 R^2 $\overline{\mathbf{R}^6}$ $\overline{\mathbf{R}^1}$ R³ $\overline{\mathbf{X}}$ Base CH₃ 2,8-difluoro-H H O monophosphate adenine CH₃ adenine monophosphate H H 0 2-(N,N-diacetyl)monophosphate $\overline{\mathbf{H}}$ H CH_3 S guanine H H CH₃ S 6-O-acetyl monophosphate guanine 8-fluoroguanine H CH₃ S monophosphate Η H CH₃ S guanine Н monophosphate 6-(N,N-diacetyl)-H CH₃ S monophosphate H adenine CH₃ 2-fluoroadenine monophosphate Η H S Η H CH₃ S 8-fluoroadenine monophosphate CH₃ 2,8-difluoro-Н S monophosphate Η adenine CH₃ adenine H Η S monophosphate CH₃ H 0 2-(N,N-diacetyl)-H diphosphate guanine 6-O-acetyl diphosphate Н Н CH_3 0 guanine CH₃ 8-fluoroguanine diphosphate H Η 0 CH₃ diphosphate H H O guanine H Η CH_3 0 6-(N,N-diacetyl)diphosphate adenine CH₃ 2-fluoroadenine diphosphate H H O CH₃ 8-fluoroadenine H Η O diphosphate 2,8-difluoro-H Η CH₃ O diphosphate adenine H CH_3 adenine diphosphate H O CH₃ 2-(N,N-diacetyl)diphosphate H H S guanine

| WO 01/90121 R ¹ | \mathbb{R}^2 | \mathbb{R}^3 | R ⁶ | X | PCT/US01/16671 Base |
|-------------------------------|----------------|----------------|-----------------|---|------------------------|
| diphosphate | H | H | CH ₃ | S | 6-O-acetyl |
| - | | | | | guanine |
| diphosphate | H | Н | CH ₃ | S | 8-fluoroguanine |
| diphosphate | H | H | CH ₃ | S | guanine |
| diphosphate | H | Н | CH ₃ | S | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| diphosphate | Н | H | CH ₃ | S | 2-fluoroadenine |
| diphosphate | Н | H | CH ₃ | S | 8-fluoroadenine |
| diphosphate | H | Н | CH ₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| diphosphate | H | H | CH ₃ | S | adenine |
| triphosphate | H | Н | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| triphosphate | H | H | CH ₃ | 0 | 6-O-acetyl |
| | | | | | guanine |
| triphosphate | Н | Н | CH ₃ | 0 | 8-fluoroguanine |
| triphosphate | Н | H | CH ₃ | 0 | guanine |
| triphosphate | H | Н | CH ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| triphosphate | Н | Н | CH ₃ | 0 | 2-fluoroadenine |
| triphosphate | Н | Н | CH ₃ | 0 | 8-fluoroadenine |
| triphosphate | Н | Н | CH ₃ | 0 | 2,8-difluoro- |
| | | 000 | | | adenine |
| triphosphate | Н | Н | CH ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| triphosphate | H | Н | CH ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| triphosphate | H | Н | CH ₃ | S | 8-fluoroguanine |
| triphosphate | H | Н | CH ₃ | S | guanine |
| triphosphate | H | Н | CH ₃ | S | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| triphosphate | H | . Н | CH ₃ | S | 2-fluoroadenine |

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|----------------------------|----------------|----------------|-----------------|---|-------------------|
| \mathbb{R}^1 | \mathbb{R}^2 | \mathbb{R}^3 | R ⁶ | X | Base |
| triphosphate | Н | Н | CH ₃ | S | 8-fluoroadenine |
| triphosphate | Н | Н | CH ₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| triphosphate | Н | H | CH ₃ | S | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2-(N,N-diacetyl)- |
| | | | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 6-O-acetyl |
| | | | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroguanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 8-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | 0 | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2-(N,N-diacetyl)- |
| | | ļ | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 6-O-acetyl |
| | | | | | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 8-fluoroguanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | guanine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 6-(N,N-diacetyl)- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 8-fluoroadenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | 2,8-difluoro- |
| | | | | | adenine |
| monophosphate | monophosphate | monophosphate | CF ₃ | S | adenine |
| acetyl | acetyl | acetyl | CF ₃ | 0 | guanine |
| acetyl | acetyl | acetyl | CF ₃ | S | guanine |

| \mathbb{R}^{1} | R ² | R ³ | R ⁶ | X | Base |
|------------------|----------------|----------------|-------------------|---|---------|
| acetyl | acetyl | acetyl | 2-bromo- vinyl | 0 | guanine |
| acetyl | acetyl | acetyl | 2-bromo- vinyl | S | guanine |

Alternatively, the following nucleosides of Formula XIV are prepared, using the appropriate sugar and pyrimidine or purine bases.

$$R^{1}O$$
 R^{6}
 X
 OR^{2}
 (XIV)

wherein:

| R ¹ | R ² | R ⁶ | X | Base |
|----------------|----------------|-----------------|---|---------------------------|
| Н | H | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| Н | H | CH ₃ | 0 | Hypoxanthine |
| H | Н | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| H | H | CH ₃ | 0 | Thymine |
| Н | Н | CH ₃ | 0 | Cytosine |
| H | Н | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| Н | Н | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| H | Н | CH ₃ | 0 | Uracil |
| H | H | CH ₃ | 0 | 5-Fluorouracil |
| H | Н | CH ₃ | S | 2,4-O-Diacetyluracil |
| H | Н | CH ₃ | S | Hypoxanthine |
| Н | H | CH ₃ | S | 2,4-O-Diacetylthymine |
| H | Н | CH ₃ | S | Thymine |
| H | Н | CH ₃ | S | Cytosine |
| H | H | CH ₃ | S | 4-(N-mono-acetyl)cytosin |

| \mathbb{R}^1 | \mathbb{R}^2 | R ⁶ | X | Base |
|----------------|----------------|-----------------|---|---------------------------|
| H | H | ĊH ₃ | S | 4-(N,N-diacetyl)cytosine |
| H | Н | CH ₃ | S | Uracil |
| H | H | CH ₃ | S | 5-Fluorouracil |
| monophosphate | H | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | H | CH ₃ | 0 | Hypoxanthine |
| monophosphate | H | CH ₃ | 0 | 2,4-O-Diacetylthym |
| monophosphate | Н | CH ₃ | 0 | Thymine |
| monophosphate | H | CH ₃ | 0 | Cytosine |
| monophosphate | Н | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytos |
| monophosphate | Н | CH ₃ | 0 | Uracil |
| monophosphate | Н | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | Н | CH ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | Н | CH ₃ | S | Hypoxanthine |
| monophosphate | Н | CH ₃ | S | 2,4-O-Diacetylthym |
| monophosphate | Н | CH ₃ | S | Thymine |
| monophosphate | Н | CH ₃ | S | Cytosine |
| monophosphate | Н | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | Н | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | Н | CH ₃ | S | Uracil |
| monophosphate | Н | CH ₃ | S | 5-Fluorouracil |
| diphosphate | Н | CH ₃ | О | 2,4-O-Diacetyluracil |
| diphosphate | Н | CH ₃ | 0 | Hypoxanthine |
| diphosphate | H | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| diphosphate | Н | CH ₃ | 0 | Thymine |
| diphosphate | Н | CH ₃ | 0 | Cytosine |
| diphosphate | Н | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| diphosphate | H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| diphosphate | Н | CH ₃ | 0 | Uracil |
| diphosphate | Н | CH ₃ | О | 5-Fluorouracil |
| diphosphate | H | CH ₃ | S | 2,4-O-Diacetyluracil |

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|---------------|----------------|-----------------|---|---------------------------|
| R' | R ² | R ⁶ | X | Base |
| diphosphate | Н | CH ₃ | S | Hypoxanthine |
| diphosphate | Н | CH ₃ | S | 2,4-O-Diacetylthymine |
| diphosphate | Н | CH ₃ | S | Thymine |
| diphosphate | Н | CH ₃ | S | Cytosine |
| triphosphate | Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| triphosphate | Н | CH ₃ | 0 | Hypoxanthine |
| triphosphate | Н | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| triphosphate | H | CH ₃ | 0 | Thymine |
| triphosphate | Н | CH ₃ | 0 | Cytosine |
| triphosphate | H | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| triphosphate | Н | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| triphosphate | Н | CH ₃ | 0 | Uracil |
| triphosphate | H | CH ₃ | 0 | 5-Fluorouracil |
| triphosphate | H | CH ₃ | S | 2,4-O-Diacetyluracil |
| triphosphate | H | CH ₃ | S | Hypoxanthine |
| triphosphate | Н | CH ₃ | S | 2,4-O-Diacetylthymine |
| triphosphate | H | CH ₃ | S | Thymine |
| triphosphate | Н | CH ₃ | S | Cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | monophosphate | CF ₃ | 0 | Hypoxanthine |
| monophosphate | monophosphate | CF ₃ | 0 | 2,4-O-Diacetylthymine |
| monophosphate | monophosphate | CF ₃ | 0 | Thymine |
| monophosphate | monophosphate | CF ₃ | 0 | Cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | 0 | Uracil |
| monophosphate | monophosphate | CF ₃ | 0 | 5-Fluorouracil |
| monophosphate | monophosphate | CF ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | monophosphate | CF ₃ | S | Hypoxanthine |
| monophosphate | monophosphate | CF ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | monophosphate | CF ₃ | S | Thymine |

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|------------------|----------------|-----------------|---|---------------------------|
| \mathbb{R}^{1} | R ² | R ⁶ | X | Base |
| monophosphate | monophosphate | CF ₃ | S | Cytosine |
| monophosphate | monophosphate | CF ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | monophosphate | CF ₃ | S | Uracil |
| monophosphate | monophosphate | CF ₃ | S | 5-Fluorouracil |
| acetyl . | acetyl | CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| acetyl | acetyl | CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| acetyl | acetyl | 2-bromo- | 0 | 4-(N,N-diacetyl)cytosine |
| | | vinyl | | |
| acetyl | acetyl | 2-bromo- | S | 4-(N,N-diacetyl)cytosine |
| | | vinyl | | |

Alternatively, the following nucleosides of Formula XV are prepared, using the appropriate sugar and pyrimidine or purine bases.

wherein:

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| R ¹ | R ⁶ | X | Base |
|----------------|-----------------|---|---------------------------|
| Н | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| H | CH ₃ | 0 | Hypoxanthine |
| H | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| H | CH ₃ | 0 | Thymine |
| H | CH ₃ | 0 | Cytosine |
| H | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| H | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| H | CH ₃ | 0 | Uracil |

| WO 01/90121 | R ⁶ | X | Base PCT/ |
|---------------|-----------------|---|---------------------------|
| <u></u> | CH ₃ | 0 | 5-Fluorouracil |
| H | | | 2,4-O-Diacetyluracil |
| H | CH ₃ | S | |
| H | CH₃ | S | Hypoxanthine |
| H | CH ₃ | S | 2,4-O-Diacetylthymine |
| H | CH ₃ | S | Thymine |
| Н | CH ₃ | S | Cytosine |
| Н | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| Н | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| Н | CH ₃ | S | Uracil |
| H | CH ₃ | S | 5-Fluorouracil |
| monophosphate | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| monophosphate | CH ₃ | 0 | Hypoxanthine |
| monophosphate | CH ₃ | 0 | 2,4-O-Diacetylthymine |
| monophosphate | CH ₃ | 0 | Thymine |
| monophosphate | CH ₃ | 0 | Cytosine |
| monophosphate | CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| monophosphate | CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| monophosphate | CH ₃ | 0 | Uracil |
| monophosphate | CH ₃ | 0 | 5-Fluorouracil |
| monophosphate | CH ₃ | S | 2,4-O-Diacetyluracil |
| monophosphate | CH ₃ | S | Hypoxanthine |
| monophosphate | CH ₃ | S | 2,4-O-Diacetylthymine |
| monophosphate | CH ₃ | S | Thymine |
| monophosphate | CH ₃ | S | Cytosine |
| monophosphate | CH ₃ | S | 4-(N-mono-acetyl)cytosine |
| monophosphate | CH ₃ | S | 4-(N,N-diacetyl)cytosine |
| monophosphate | CH ₃ | S | Uracil |
| monophosphate | CH ₃ | S | 5-Fluorouracil |
| diphosphate | CH ₃ | 0 | 2,4-O-Diacetyluracil |
| diphosphate | CH ₃ | 0 | Hypoxanthine |
| diphosphate | CH ₃ | 0 | 2,4-O-Diacetylthymine |

| R ⁶ | X | Base |
|-----------------|---|---|
| CH ₃ | | |
| | 0 | Thymine |
| CH ₃ | 0 | Cytosine |
| CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| CH ₃ | 0 | Uracil |
| CH ₃ | 0 | 5-Fluorouracil |
| CH ₃ | S | 2,4-O-Diacetyluracil |
| CH ₃ | S | Hypoxanthine |
| CH ₃ | S | 2,4-O-Diacetylthymine |
| CH ₃ | S | Thymine |
| CH₃ | S | Cytosine |
| CH ₃ | 0 | 2,4-O-Diacetyluracil |
| CH ₃ | 0 | Hypoxanthine |
| CH ₃ | 0 | 2,4-O-Diacetylthymine |
| CH ₃ | 0 | Thymine |
| CH ₃ | 0 | Cytosine |
| CH ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| CH ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| CH ₃ | 0 | Uracil |
| CH ₃ | 0 | 5-Fluorouracil |
| CH ₃ | S | 2,4-O-Diacetyluracil |
| CH ₃ | S | Hypoxanthine |
| CH ₃ | S | 2,4-O-Diacetylthymine |
| CH ₃ | S | Thymine |
| CH ₃ | S | Cytosine |
| CF ₃ | 0 | 2,4-O-Diacetyluracil |
| CF ₃ | 0 | Hypoxanthine |
| CF ₃ | 0 | 2,4-O-Diacetylthymine |
| CF ₃ | 0 | Thymine |
| CF ₃ | 0 | Cytosine |
| CF ₃ | 0 | 4-(N-mono-acetyl)cytosine |
| | CH ₃ | CH3 O CH3 O CH3 O CH3 O CH3 S CH3 S CH3 S CH3 S CH3 O CH3 S CH4 S CH4 S CH4 S CH4 S |

| | PCT/ | |
|-----------------|---|---|
| \mathbb{R}^6 | X | Base |
| CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| CF ₃ | 0 | Uracil |
| CF ₃ | 0 | 5-Fluorouracil |
| CF ₃ | S | 2,4-O-Diacetyluracil |
| CF ₃ | S | Hypoxanthine |
| CF ₃ | S | 2,4-O-Diacetylthymine |
| CF ₃ | S | Thymine |
| CF ₃ | S | Cytosine |
| CF ₃ | S | 4-(N-mono-acetyl)cytosine |
| CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| CF ₃ | S | Uracil |
| CF ₃ | S | 5-Fluorouracil |
| CF ₃ | 0 | 4-(N,N-diacetyl)cytosine |
| CF ₃ | S | 4-(N,N-diacetyl)cytosine |
| 2-bromo-vinyl | 0 | 4-(N,N-diacetyl)cytosine |
| 2-bromo-vinyl | S | 4-(N,N-diacetyl)cytosine |
| | CF ₃ | CF3 O CF3 O CF3 O CF3 S 2-bromo-vinyl O |

Alternatively, the following nucleosides of Formula XVIII are prepared, using the appropriate sugar and pyrimidine or purine bases.

wherein:

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| R ¹ | R ⁶ | \mathbb{R}^7 | X | Base | R ⁸ | R ⁹ |
|----------------|-----------------|----------------|---|-----------------------|----------------|----------------|
| H | CH ₃ | OH | 0 | 2,4-O-Diacetyluracil | H | Me |
| H | CH ₃ | OH | 0 | Hypoxanthine | H | Me |
| H | CH ₃ | OH | 0 | 2,4-O-Diacetylthymine | H | Me |

| WO 01/90121 R ¹ | R ⁶ | R ⁷ | X | Base | /US01/10 R ⁸ | R ⁹ |
|-------------------------------|-----------------|----------------|---|---------------------------|----------------------------|----------------|
| H | CH ₃ | OH | 0 | Thymine | H | Me |
| H | CH ₃ | OH | 0 | Cytosine | Н | Me |
| H | CH ₃ | OH | 0 | 4-(N-mono-acetyl)cytosine | Н | Me |
| H . | CH ₃ | OH | 0 | 4-(N,N-diacetyl)cytosine | H | Me |
| H | CH ₃ | OH | 0 | Uracil | Н | Me |
| Н | CH ₃ | OH | 0 | 5-Fluorouracil | H | Me |
| H | CH ₃ | OH | S | 2,4-O-Diacetyluracil | H | Me |
| H | CH ₃ | OH | S | Hypoxanthine | Н | Me |
| H | CH ₃ | OH | S | 2,4-O-Diacetylthymine | H | Me |
| Н | CH ₃ | OH | S | Thymine | H | Me |
| H | CH ₃ | OH | S | Cytosine | Н | Me |
| H | CH ₃ | OH | S | 4-(N-mono-acetyl)cytosine | Н | Me |
| H | CH ₃ | OH | S | 4-(N,N-diacetyl)cytosine | Н | Me |
| H | CH ₃ | OH | S | Uracil | Н | Me |
| H | CH ₃ | ОН | S | 5-Fluorouracil | Н | Me |
| monophosphate | CH ₃ | OH | 0 | 2,4-O-Diacetyluracil | H | Me |
| monophosphate | CH ₃ | OH | 0 | Hypoxanthine | Н | Me |
| monophosphate | CH ₃ | OH | 0 | 2,4-O-Diacetylthymine | H | Me |
| monophosphate | CH ₃ | OH | 0 | Thymine | H | Me |
| monophosphate | CH ₃ | OH | 0 | Cytosine | H | Me |
| monophosphate | CH ₃ | OH | 0 | 4-(N-mono-acetyl)cytosine | Н | Me |
| monophosphate | CH ₃ | OH | 0 | 4-(N,N-diacetyl)cytosine | Н | Me |
| monophosphate | CH ₃ | OH | 0 | Uracil | Н | Me |
| monophosphate | CH ₃ | OH | 0 | 5-Fluorouracil | H | Me |
| monophosphate | CH ₃ | OH | S | 2,4-O-Diacetyluracil | H | Me |
| monophosphate | CH ₃ | ОН | S | Hypoxanthine | Н | Me |
| monophosphate | CH ₃ | OH | S | 2,4-O-Diacetylthymine | Н | Me |
| monophosphate | CH ₃ | OH | S | Thymine | H | Me |
| monophosphate | CH ₃ | ОН | S | Cytosine | H | Me |
| monophosphate | CH ₃ | OH | S | 4-(N-mono-acetyl)cytosine | H | Me |
| monophosphate | CH ₃ | OH | S | 4-(N,N-diacetyl)cytosine | Н | Me |

| monophosphate Comonophosphate | R ⁶ CH ₃ CH ₃ CH ₃ CH ₃ | OH OH | X S | Base Uracil | R ⁸ | R ⁹ |
|--|--|-------|--------|---------------------------|----------------|----------------|
| monophosphate Codiphosphate Co | CH ₃ | OH | | Uracil | Н | Me |
| diphosphate Codiphosphate Codi | CH ₃ | | S | | | |
| diphosphate Codiphosphate Codi | | OT. | | 5-Fluorouracil | H | Me |
| diphosphate C | CH ₃ | OH | 0 | 2,4-O-Diacetyluracil | H | Me |
| diphosphate C | 1 | OH | 0 | Hypoxanthine | H | Me |
| diphosphate C di | CH ₃ | OH | 0 | 2,4-O-Diacetylthymine | Н | Me |
| diphosphate C | CH ₃ | OH | 0 | Thymine | Н | Me |
| diphosphate C | CH ₃ | OH | 0 | Cytosine | Н | Me |
| diphosphate C diphosphate C diphosphate C diphosphate C diphosphate C | CH ₃ | OH | 0 | 4-(N-mono-acetyl)cytosine | Н | Me |
| diphosphate C diphosphate C diphosphate C | CH ₃ | OH | 0 | 4-(N,N-diacetyl)cytosine | H | Me |
| diphosphate C | CH ₃ | OH | 0 | Uracil | Н | Me |
| diphosphate C | CH ₃ | OH | 0 | 5-Fluorouracil | Н | Me |
| | CH ₃ | OH | S | 2,4-O-Diacetyluracil | Н | Me |
| diphosphate C | CH ₃ | OH | S | Hypoxanthine | Н | Me |
| | CH ₃ | OH | S | 2,4-O-Diacetylthymine | Н | Me |
| diphosphate C | CH ₃ | OH. | S | Thymine | H | Me |
| diphosphate C | CH ₃ | ОН | S | Cytosine | Н | Me |
| triphosphate C | CH ₃ | ОН | 0 | 2,4-O-Diacetyluracil | Н | Me |
| triphosphate C | CH ₃ | ОН | 0 | Hypoxanthine | Н | Me |
| triphosphate C | CH₃ | ОН | 0 | 2,4-O-Diacetylthymine | H | Me |
| triphosphate C | CH ₃ | ОН | 0 | Thymine | Н | Me |
| triphosphate C | CH ₃ | OH | 0 | Cytosine | Н | Me |
| triphosphate C | CH ₃ | OH | 0 | 4-(N-mono-acetyl)cytosine | H | Me |
| triphosphate C | CH ₃ | ОН | 0 | 4-(N,N-diacetyl)cytosine | H | Me |
| triphosphate C | CH ₃ | OH | 0 | Uracil | Н | Me |
| triphosphate C | CH ₃ | OH | 0 | 5-Fluorouracil | Н | Me |
| triphosphate C | CH ₃ | OH | S | 2,4-O-Diacetyluracil | H | Me |
| triphosphate C | CH ₃ | ОН | S | Hypoxanthine | Н | Me |
| triphosphate C | CH ₃ | OH | S | 2,4-O-Diacetylthymine | Н | Me |
| triphosphate C | CH ₃ | OH | S | Thymine | Н | Me |
| triphosphate C | CH ₃ | OH | S | Cytosine | Н | Me |
| monophosphate C | | OH | 0 | 2,4-O-Diacetyluracil | Н | Me |

| R ¹ | R ⁶ | \mathbf{R}^7 | X | Base | R ⁸ | R ⁹ |
|----------------|-----------------|----------------|---|---------------------------|----------------|----------------|
| monophosphate | CF ₃ | OH | 0 | Hypoxanthine | H | Me |
| monophosphate | CF ₃ | OH | 0 | 2,4-O-Diacetylthymine | H | Me |
| monophosphate | CF ₃ | OH | 0 | Thymine | Н | Me |
| monophosphate | CF ₃ | OH | 0 | Cytosine | H | Me |
| monophosphate | CF ₃ | ОН | 0 | 4-(N-mono-acetyl)cytosine | Н | Me |
| monophosphate | CF ₃ | ОН | 0 | 4-(N,N-diacetyl)cytosine | H | Me |
| monophosphate | CF ₃ | ОН | 0 | Uracil | H | Me |
| monophosphate | CF ₃ | ОН | 0 | 5-Fluorouracil | H | Me |
| monophosphate | CF ₃ | OH | S | 2,4-O-Diacetyluracil | H | Me |
| monophosphate | CF ₃ | ОН | S | Hypoxanthine | H | Me |
| monophosphate | CF ₃ | OH | S | 2,4-O-Diacetylthymine | H | Me |
| monophosphate | CF ₃ | OH | S | Thymine | H | Me |
| monophosphate | CF ₃ | OH | S | Cytosine | H | Me |

PCT/US01/16671

Н

Η

H

Η

Η

H

Me

Me

Me

Me

Br

Br

VII. Anti-Hepatitis C Activity

CF₃

CF₃

CF₃

 CH_3

 CH_3

monophosphate

monophosphate

monophosphate

monophosphate

acetyl

acetyl

5

10

OH

OH

OH

OH

OH

OH

S

S

S

S

0

S

Uracil

5-Fluorouracil

4-(N-mono-acetyl)cytosine

4-(N,N-diacetyl)cytosine

4-(N,N-diacetyl)cytosine

4-(N,N-diacetyl)cytosine

WO 01/90121

Compounds can exhibit anti-hepatitis C activity by inhibiting HCV polymerase, by inhibiting other enzymes needed in the replication cycle, or by other pathways. A number of assays have been published to assess these activities. A general method that assesses the gross increase of HCV virus in culture is disclosed in U.S. Patent No. 5,738,985 to Miles et al. In vitro assays have been reported in Ferrari et al., Inl. of Vir., 73:1649-1654, 1999; Ishii et al., Hepatology, 29:1227-1235,1999; Lohmann et al., Inl. of Bio. Chem., 274:10807-10815, 1999; and Yamashita et al., Inl. of Bio. Chem., 273:15479-15486, 1998.

WO 97/12033, filed on September 27, 1996, by Emory University, listing C. Hagedorn and A. Reinoldus as inventors, and which claims priority to U.S.S.N. 60/004,383,

filed on September 1995, describes an HCV polymerase assay that can be used to evaluate the activity of the compounds described herein. Another HCV polymerase assay has been reported by Bartholomeusz, et al., Hepatitis C virus (HCV) RNA polymerase assay using cloned HCV non-structural proteins; Antiviral Therapy 1996:1(Supp 4) 18-24.

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Screens that measure reductions in kinase activity from HCV drugs are disclosed in U.S. Patent No. 6,030,785, to Katze *et al.*, U.S. Patent No. 6,010,848 to Delvecchio et al, and U.S. Patent No. 5,759,795 to Jubin *et al.* Screens that measure the protease inhibiting activity of proposed HCV drugs are disclosed in U.S. Patent No. 5,861,267 to Su et al, U.S. Patent No. 5,739,002 to De Francesco et al, and U.S. Patent No. 5,597,691 to Houghton *et al*.

Example 4: Phosphorylation Assay of Nucleoside to Active Triphosphate

To determine the cellular metabolism of the compounds, HepG2 cells were obtained from the American Type Culture Collection (Rockville, MD), and were grown in 225 cm² tissue culture flasks in minimal essential medium supplemented with non-essential amino acids, 1% penicillin-streptomycin. The medium was renewed every three days, and the cells were subcultured once a week. After detachment of the adherent monolayer with a 10 minute exposure to 30 mL of trypsin-EDTA and three consecutive washes with medium, confluent HepG2 cells were seeded at a density of 2.5×10^6 cells per well in a 6-well plate and exposed to $10 \mu M$ of [3H] labeled active compound (500 dpm/pmol) for the specified time periods. The cells were maintained at $37^{\circ}C$ under a 5% CO₂ atmosphere. At the selected time points, the cells were washed three times with ice-cold phosphate-buffered saline (PBS). Intracellular active compound and its respective metabolites were extracted by incubating the cell pellet overnight at $-20^{\circ}C$ with 60% methanol followed by extraction with an additional $20 \mu L$ of cold methanol for one hour in an ice bath. The extracts were then combined, dried under gentle filtered air flow and stored at $-20^{\circ}C$ until HPLC analysis. The preliminary results of the HPLC analysis are tabulated in Table 1.

Table 1

| | [pmol/million cells] | | | | | | |
|----------|--------------------------------------|--------------------------------------|--------------------------------------|--------------------------------------|--|--|--|
| Time (h) | β-D-2'-CH ₃ - riboA-TP | β-D-2'-CH ₃ - riboU-TP | β-D-2'-CH ₃ - riboC-TP | β-D-2'-CH ₃ - riboG-TP | | | |
| 2 | 33.1 | 0.40 | 2.24 | ND | | | |
| 4 | 67.7 | 1.21 | 3.99 | ND | | | |
| 8 | 147 | 1.57 | 9.76 | 2.85 | | | |
| 24 | 427 | 6.39 | 34.9 | 0.91 | | | |
| 30 | 456 | 7.18 | 36.2 | 3.22 | | | |
| 48 | 288 | 9.42 | 56.4 | 6.26 | | | |

Example 5: Bioavailability Assay in Cynomolgus Monkeys

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Within 1 week prior to the study initiation, the cynomolgus monkey was surgically implanted with a chronic venous catheter and subcutaneous venous access port (VAP) to facilitate blood collection and underwent a physical examination including hematology and serum chemistry evaluations and the body weight was recorded. Each monkey (six total), received approximately 250 uCi of ³H activity with each dose of active compound, namely β-D-2'-CH₃-riboG at a dose level of 10 mg/kg at a dose concentration of 5 mg/mL, either via an intravenous bolus (3 monkeys, IV), or via oral gavage (3 monkeys, PO). Each dosing syringe was weighed before dosing to gravimetrically determine the quantity of formulation administered. Urine samples were collected via pan catch at the designated intervals (approximately 18-0 hours pre-dose, 0-4, 4-8 and 8-12 hours post-dosage) and processed. Blood samples were collected as well (pre-dose, 0.25, 0.5, 1, 2, 3, 6, 8, 12 and 24 hours post-dosage) via the chronic venous catheter and VAP or from a peripheral vessel if the chronic venous catheter procedure should not be possible. The blood and urine samples were analyzed for the maximum concentration (C_{max}), time when the maximum concentration was achieved (Tmax), area under the curve (AUC), half life of the dosage concentration (T₁₆), clearance (CL), steady state volume and distribution (V_{ss}) and bioavailability (F), which are tabulated in Tables 2 and 3, and graphically illustrated in Figures 2 and 3, respectively.

38.8

| | Dose (mg) | AUC (ng/mL x h) | Norm AUC (ng/mL x h/mg) | Mean Norm AUC (ng/mL x h/mg) | F (%) |
|-------------|--------------|--------------------|----------------------------|------------------------------|-------|
| IV Monkey 1 | 46.44 | 13614 | 293.2 | | |
| IV Monkey 2 | 24.53 | 6581 | 268.3 | · | |
| IV Monkey 3 | 20.72 | 6079 | 293.4 | 284.9 | |
| PO Monkey 1 | 29.04 | 758 | 26.1 | | |
| 2011 | 20.02 | 000 | 20.0 | | |

Table 3: Experimental Pharmacokinetics of β-D-2'-CH₃-riboG in Cynomolgus Monkeys

| | IV | PO | | |
|--------------------------|---------------------|--------------------|--|--|
| Dose/Route (mg/kg) | 10 | 10 | | |
| C _{max} (ng/mL) | 6945.6 ± 1886.0 | 217.7 ± 132.1 | | |
| T _{max} (hr) | 0.25 ± 0.00 | 2.00 ± 1.00 | | |
| AUC (ng/mL x hr) | 8758.0 ± 4212.9 | 1166.0 ± 589.6 | | |
| T _{1/2} (hr) | 7.9 ± 5.4 | 10.3 ± 4.1 | | |
| CL (L/hr/kg) | 1.28 ± 0.48 | | | |
| V _{ss} (L/kg) | 2.09 ± 0.54 | | | |
| F (%) | 13.8 | | | |

Example 6: Bone Marrow Toxicity Assay

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Human bone marrow cells were collected from normal healthy volunteers and the mononuclear population was separated by Ficoll-Hypaque gradient centrifugation as described previously by Sommadossi J-P, Carlisle R. "Toxicity of 3'-azido-3'deoxythymidine and 9-(1,3-dihydroxy-2-propoxymethyl)guanine for normal human hematopoietic progenitor cells in vitro" Antimicrobial Agents and Chemotherapy 1987; 31:452-454; and Sommadossi J-P, Schinazi RF, Chu CK, Xie M-Y. "Comparison of cytotoxicity of the (-)- and (+)-enantiomer of 2',3'-dideoxy-3'-thiacytidine in normal human bone marrow progenitor cells" Biochemical Pharmacology 1992; 44:1921-1925. The culture assays for CFU-GM and BFU-E were performed using a bilayer soft agar or methylcellulose method. Drugs were diluted in tissue culture medium and filtered. After 14 to 18 days at 37°C in a humidified atmosphere of 5% CO₂ in air, colonies of greater than 50 cells were counted using an inverted microscope. The results in Table 4 are presented as the percent inhibition of colony formation in the presence of drug compared to solvent control cultures.

Table 4: Human Bone Marrow Toxicity CFU-GM and BFU-E Clonogenic Assays

| : | IC ₅₀ in μM | |
|-------------------------------|------------------------|-------|
| Treatment | CFU-GM | BFU-E |
| ribavirin | ~ 5 | ~ 1 |
| β-D-2'-CH ₃ -riboA | > 100 | > 100 |
| β-D-2'-CH ₃ -riboU | > 100 | > 100 |
| β-D-2'-CH ₃ -riboC | > 10 | > 10 |
| β-D-2'-CH ₃ -riboG | > 10 | > 100 |

Example 7: Mitochondria Toxicity Assay

HepG2 cells were cultured in 12-well plates as described above and exposed to various concentrations of drugs as taught by Pan-Zhou X-R, Cui L, Zhou X-J, Sommadossi J-P, Darley-Usmer VM. "Differential effects of antiretroviral nucleoside analogs on mitochondrial function in HepG2 cells" Antimicrob Agents Chemother 2000; 44:496-503. Lactic acid levels in the culture medium after 4 day drug exposure was measured using a Boehringer lactic acid assay kit. Lactic acid levels were normalized by cell number as measured by hemocytometer count. The preliminary results from this assay are tabulated in Table 5.

Table 5: Mitochondrial Toxicity Study (L-lactic acid assay)

| 1 | Conc. (µM) | lactate (mg/10 ⁶ cell) | % of Control |
|-------------------------------|------------|-----------------------------------|--------------|
| Control | | 2.18 | |
| FIAU | 10 | 3.73 | 170.4 |
| β-D-2'-CH ₃ -riboC | 1 | 2.52 | 115.3 |
| | 10 | 2.36 | 107.9 |
| | 50 | 2.26 | 103.4 |
| | 100 | 2.21 | 101.2 |

FIAU

β-D-2'-CH₃-riboC

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This invention has been described with reference to its preferred embodiments. Variations and modifications of the invention, will be obvious to those skilled in the art from the foregoing detailed description of the invention.

We Claim:

1. A compound of Formula I:

$$X^1$$
 X^1
 X^2
 CH_3
 CH_3
 (I)

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

2. A compound of Formula II:

$$X^1$$
 N
 N
 X^2
 $OR^2 \cdot OR^3$
(II)

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

3. A compound of Formula III:

$$X^{1}$$
 N
 N
 X^{2}
 CH_{3}
 OR^{2}
 OR^{3}
(III)

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is

capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR4, NR4R5 or SR4;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

4. A compound of Formula IV:

or a pharmaceutically acceptable salt thereof, wherein:

 R^1 , R^2 and R^3 are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 and R^3 are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

5. A compound of Formula V:

$$X^1$$
 X^1
 X^1

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

6. A compound of Formula VI:

or a pharmaceutically acceptable salt thereof, wherein:

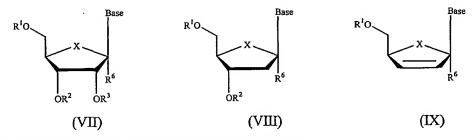
R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

7. A compound selected from Formulas VII, VIII and IX:



or a pharmaceutically acceptable salt thereof, wherein:

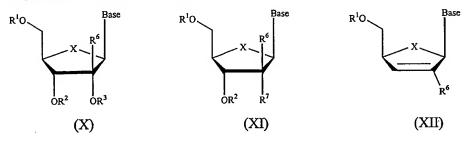
Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is

capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂, or CH₂.

8. A compound of Formulas X, XI and XII:



or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and

9. A compound selected from Formulas XIII, XIV and XV:

R¹⁰
Base
$$R^6$$
 X
 OR^2
 OR^3
 OR^2
 OR^2
 OR^3
 OR^2
 OR^3
 OR^2
 OR^3
 OR^2
 OR^3
 OR^2
 OR^3
 OR^3
 OR^2
 OR^3
 OR^3
 OR^3
 OR^3
 OR^3
 OR^3

or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(acyl)₂; and X is O, S, SO₂ or CH₂.

10. A compound of Formula XVI:

$$R^{10}$$
 R^{10} R^{8} R^{6} R^{7} R^{8} R^{7}

or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

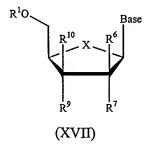
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine;

alternatively, R^7 and R^9 , R^7 and R^{10} , R^8 and R^9 , or R^8 and R^{10} can come together to form a bond; and

X is O, S, SO₂ or CH₂.

11. A compound of Formula XVII:



or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

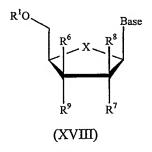
R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl

(including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -O(alkyl), -N(lower alkyl), -N(lower alkyl), -N(lower alkyl), -N(acyl);

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a bond; and X is O, S, SO₂ or CH₂.

12. A compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or

other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkyl), -O(alkyl), -N(lower alkyl), -N(lower alkyl), -N(lower alkyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

 R^8 is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R^7 and R^9 , or R^8 and R^9 can come together to form a bond; and X is O, S, SO₂ or CH₂.

13. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

14. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

15. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

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16. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

17. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

18. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

19. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

20. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

21. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

22. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

23. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

24. A compound of the structure:

or a pharmaceutically acceptable salt thereof.

- 25. The compound as described in any of the preceding claims 1-24, wherein the said compound is in the form of a dosage unit.
- 26. The compound as described in claim 187, wherein the dosage unit contains 10 to 1500 mg of said compound.
- 27. The compound as described in claim 187 or 188, wherein said dosage unit is a tablet or capsule.
- 28. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula I:

$$X^1$$
 X^1
 X^2
 CH_3
 CH_3
 (I)

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more

substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

29. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula II:

$$X^{1}$$
 N
 N
 X^{2}
 N
 N
 X^{2}
 N
 N
 X^{2}
 N
 N
 X
 X

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

30. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula III:

$$X^1$$
 N
 N
 N
 X^2
 OR^2
 OR^3
(III)

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

31. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula IV:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

32. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula V:

$$X^1$$
 X^1
 X^1

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

33. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula VI:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

34. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formulas VII, VIII or IX:

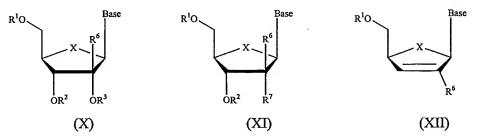
or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

35. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula X, XI or XII:



or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

Base is a purine or pyrimidine base as defined herein;

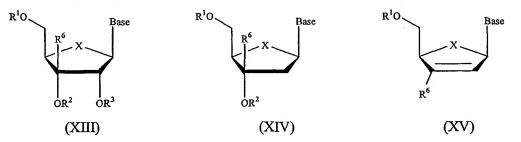
R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid,

including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

36. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XIII, XIV or XV:



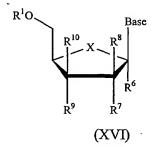
or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

37. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XVI:



or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

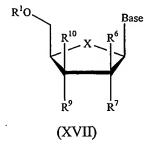
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂ or -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a bond; and

X is O, S, SO₂ or CH₂.

38. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XVII:



or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

Base is a purine or pyrimidine base as defined herein;

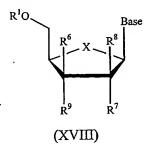
R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a bond; and X is O, S, SO₂ or CH₂.

39. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a bond; and X is O, S, SO₂ or CH₂.

40. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

41. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

42. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

43. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

44. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

45. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

46. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

47. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

48. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

49. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

50. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

51. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier or diluent.

52. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula I:

$$X^1$$
 X^1
 X^2
 CH_3
 CH_3
 CH_3

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

 R^1 , R^2 and R^3 are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 and R^3 are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

53. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula II:

$$X^1$$
 N
 N
 X^2
 N
 N
 X^2
 N
 N
 X^2
 N
 N
 N
 X^2
 N
 N
 X
 X

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

54. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula III:

$$X^1$$
 N
 N
 X^2
 OR^2
 OR^3
(III)

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

55. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula IV:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

56. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula V:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

57. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula VI:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

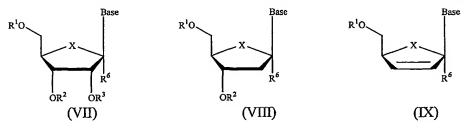
R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

58. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula VII, VIII or IX:



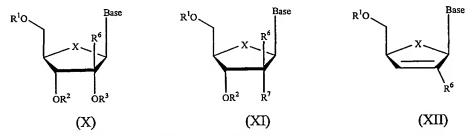
or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂, or CH₂.

59. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula X, XI or XII:



or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

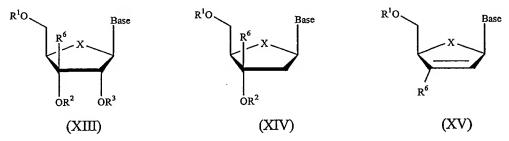
R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with

one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂, or CH₂.

60. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XIII, XIV or XV:



or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

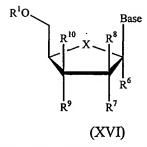
R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is

capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and

X is O, S, SO₂ or CH₂.

61. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XVI:



or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

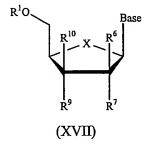
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine;

alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a bond; and

X is O, S, SO₂, or CH₂.

62. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XVII:



or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

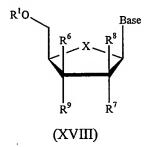
R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a bond; and X is O, S, SO₂ or CH₂.

63. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a bond; and X is O, S, SO₂ or CH₂.

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64. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

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or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

65. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

66. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

67. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

68. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

69. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

70. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

71. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

72. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

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73. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

74. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

75. A pharmaceutical composition for the treatment or prophylaxis of a Hepatitis C virus in a host, comprising an effective amount of a compound of structure:

or a pharmaceutically acceptable salt thereof, in combination with one or more other antivirally effective agents.

76. The pharmaceutical composition as described in any of the preceding claims 28-75, wherein the said compound is in the form of a dosage unit.

77. The pharmaceutical composition as described in claim 76, wherein the dosage unit contains 10 to 1500 mg of said compound.

- 78. The pharmaceutical composition as described in claim 75 or 76, wherein said dosage unit is a tablet or capsule.
- 79. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula I:

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

 X^1 and X^2 are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR^4 , NR^4NR^5 or SR^4 ; and

80. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula II:

$$X^1$$
 N
 N
 X^2
 OR^2
 OR^3
(II)

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

81. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula III:

$$X^1$$
 N
 N
 X^2
 OR^2
 OR^3
(III)

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

82. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula IV:

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

83. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula V:

$$R^{1}O$$
 $H_{3}C$
 OR^{2}
 OR^{3}
 (V)

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

84. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula VI:

or a pharmaceutically acceptable salt thereof, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

85. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula VII, VIII or IX:

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$$R^{1}O$$
 $R^{1}O$
 R

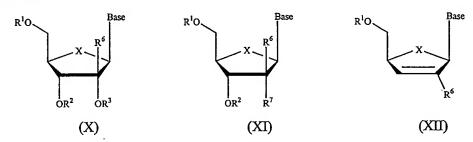
or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂, or CH₂.

86. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula X, XI or XII:



or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

87. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XIII, XIV or XV:

or a pharmaceutically acceptable salt thereof, wherein:

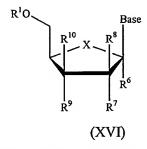
Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with

one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(acyl)₂; and X is O, S, SO₂ or CH₂.

88. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XVI:



or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

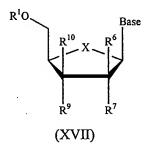
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a bond; and

X is O, S, SO₂ or CH₂.

89. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XVII:



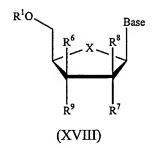
or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a bond; and X is O, S, SO₂ or CH₂.

90. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

 R^8 is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R^7 and R^9 , or R^8 and R^9 can come together to form a bond; and X is O, S, SO_2 or CH_2 .

91. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

92. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

93. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

94. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

95. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

96. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

97. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

98. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

99. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

100. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

101. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof.

102. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

103. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula I:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

104. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula II:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

105. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula III:

$$X^1$$
 N
 N
 X^2
 CH_3
 OR^2
 OR^3
(III)

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

 X^1 and X^2 are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR^4 , NR^4NR^5 or SR^4 ; and

106. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula IV:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

107. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula V:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

108. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula VI:

$$X^1$$
 X^1
 X^1

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

109. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula VII, VIII or IX:

WO 01/90121 $R^{1}O$ R^{6} R^{6} R^{6} R^{6} R^{6} R^{6} R^{6} $R^{1}O$ R^{6} R^{6} $R^{1}O$ R^{6} $R^{1}O$ R^{6} $R^{1}O$ R^{6} $R^{1}O$ R^{6} $R^{1}O$ R^{6} $R^{1}O$ R^{1}

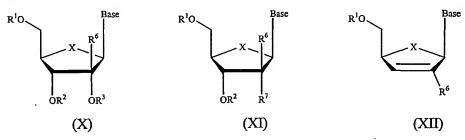
or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂, or CH₂.

110. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula X, XI or XII:



or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

111. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XIII, XIV or XV:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl

(including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

112. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XVI:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

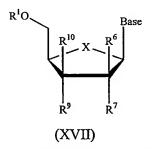
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine:

alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a bond; and

X is O, S, SO₂ or CH₂.

113. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XVII:



or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

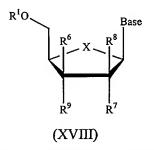
R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; R¹⁰ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁷ and R¹⁰ can come together to form a bond; and

alternatively, R' and R', or R' and R' can come together to form a bond; and X is O, S, SO₂ or CH₂.

114. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an anti-virally effective amount of a compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more other antivirally effective agents, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl

O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a bond; and X is O, S, SO₂ or CH₂.

115. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

116. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

117. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

118. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

119. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

120. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

121. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

122. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

123. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

124. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

125. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

126. A method for the treatment or prophylaxis of a Hepatitis C virus infection in a host, comprising administering an antivirally effective amount of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in combination or alternation with one or more antivirally effective agents.

- 127. Method of treatment as described in any of the preceding claims 79-126, wherein the said compound is in the form of a dosage unit.
- 128. Method of treatment as described in claim 127, wherein the dosage unit contains 10 to 1500 mg of said compound.
- 129. Method of treatment as described in claim 127 or 128, wherein said dosage unit is a tablet or capsule.
- 130. A use of a compound of Formula I:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl

and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

131. A use of a compound of Formula II:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

132. A use of a compound of Formula III:

$$X^{1}$$
 N
 X^{2}
 CH_{3}
 OR^{2}
 OR^{3}
(III)

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ and X² are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

133. A use of a compound of Formula IV:

$$X^1$$
 X^1
 X^1

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

 R^1 , R^2 and R^3 are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R^1 , R^2 and R^3 are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

134. A use of a compound of Formula V:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

135. A use of a compound of Formula VI:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid,

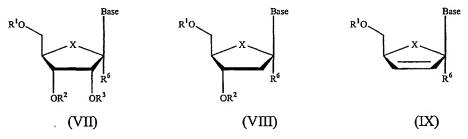
including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

136. A use of a compound selected from Formulas VII, VIII and IX:



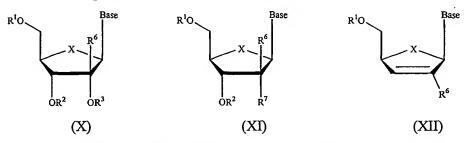
or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and

137. A use of a compound of Formulas X, XI and XII:



or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

138. A use of a compound selected from Formulas XIII, XIV and XV:

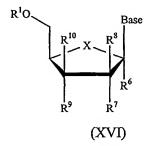
or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

139. A use of a compound of Formula XVI:



Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

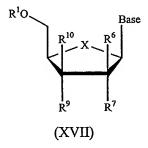
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine, or iodine;

alternatively, R^7 and R^9 , R^7 and R^{10} , R^8 and R^9 , or R^8 and R^{10} can come together to form a bond; and

X is O, S, SO_2 or CH_2 .

140. A use of a compound of Formula XVII:



or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

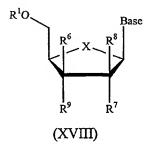
Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

 R^{10} is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; alternatively, R^7 and R^9 , or R^7 and R^{10} can come together to form a bond; and X is O, S, SO₂ or CH₂.

141. A use of a compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with

one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl)

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a bond; and X is O, S, SO₂ or CH₂.

142. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

143. A use of a compound of the structure:

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144. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

145. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

146. A use of a compound of the structure:

147. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

148. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

149. A use of a compound of the structure:

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150. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

151. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

152. A use of a compound of the structure:

153. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

154. A use of a compound of Formula I:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

 X^1 and X^2 are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR^4 , NR^4NR^5 or SR^4 ; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

155. A use of a compound of Formula II:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are is independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

 X^1 and X^2 are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR^4 , NR^4NR^5 or SR^4 ; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

156. A use of a compound of Formula III:

$$X^{1}$$
 N
 N
 N
 X^{2}
 CH_{3}
 OR^{2}
 OR^{3}
(III)

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

 X^1 and X^2 are independently selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR^4 , NR^4NR^5 or SR^4 ; and

 R^4 and R^5 are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

157. A use of a compound of Formula IV:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H, phosphate (including mono-, di- or triphosphate and a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

158. A use of a compound of Formula V:

$$X^{1}$$
 X^{1}
 X^{1

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

159. A use of a compound of Formula VI:

$$R^{1}O$$
 CH_{3}
 OR^{2}
 OR^{3}
 (VI)

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid,

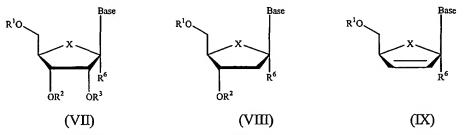
including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate; and

Y is hydrogen, bromo, chloro, fluoro, iodo, OR⁴, NR⁴R⁵ or SR⁴;

X¹ is selected from the group consisting of H, straight chained, branched or cyclic alkyl, CO-alkyl, CO-aryl, CO-alkoxyalkyl, chloro, bromo, fluoro, iodo, OR⁴, NR⁴NR⁵ or SR⁴; and

R⁴ and R⁵ are independently hydrogen, acyl (including lower acyl), or alkyl (including but not limited to methyl, ethyl, propyl and cyclopropyl).

160. A use of a compound selected from Formulas VII, VIII and IX:



or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

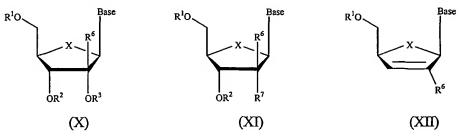
R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, 2-Br-ethyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), CF₃, chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and

X is O, S, SO₂, or CH₂.

161. A use of a compound of Formulas X, XI and XII:

Base is a purine or pyrimidine base as defined herein;



or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ is hydrogen, OR³, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

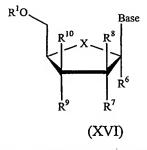
162. A use of a compound selected from Formulas XIII, XIV and XV:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein: Base is a purine or pyrimidine base as defined herein;

R¹, R² and R³ are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹, R² and R³ are independently H or phosphate;

R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂; and X is O, S, SO₂ or CH₂.

163. A use of a compound of Formula XVI:



or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

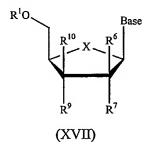
R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁸ and R¹⁰ are independently H, alkyl (including lower alkyl), chlorine, bromine or iodine;

alternatively, R⁷ and R⁹, R⁷ and R¹⁰, R⁸ and R⁹, or R⁸ and R¹⁰ can come together to form a bond; and

X is O, S, SO₂ or CH₂.

164. A use of a compound of Formula XVII:



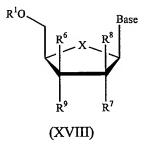
or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein: Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chloro, bromo, fluoro, iodo, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

R⁷ and R⁹ are independently hydrogen, OR², hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(alkenyl), chlorine, bromine, iodine, NO₂, NH₂, -NH(lower alkyl), -NH(acyl), -N(lower alkyl)₂, -N(acyl)₂;

 R^{10} is H, alkyl (including lower alkyl), chlorine, bromine, or iodine; alternatively, R^7 and R^9 , or R^7 and R^{10} can come together to form a bond; and X is O, S, SO₂ or CH₂.

165. A use of a compound of Formula XVIII:



or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus, wherein:

Base is a purine or pyrimidine base as defined herein;

R¹ and R² are independently H; phosphate (including monophosphate, diphosphate, triphosphate, or a stabilized phosphate prodrug); acyl (including lower acyl); alkyl (including lower alkyl); sulfonate ester including alkyl or arylalkyl sulfonyl including methanesulfonyl and benzyl, wherein the phenyl group is optionally substituted with

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one or more substituents as described in the definition of aryl given herein; a lipid, including a phospholipid; an amino acid; a carbohydrate; a peptide; a cholesterol; or other pharmaceutically acceptable leaving group which when administered *in vivo* is capable of providing a compound wherein R¹ and R² are independently H or phosphate; R⁶ is hydrogen, hydroxy, alkyl (including lower alkyl), azido, cyano, alkenyl, alkynyl, Br-vinyl, -C(O)O(alkyl), -C(O)O(lower alkyl), -O(acyl), -O(lower acyl), -O(alkyl), -O(lower alkyl), -O(alkenyl), -O(alkenyl), -N(lower alkyl), -N(lower alkyl), -N(lower alkyl), -N(lower alkyl), -N(lower alkyl), -N(lower alkyl);

R⁷ and R⁹ are independently hydrogen, OR², alkyl (including lower alkyl), alkenyl, alkynyl, Br-vinyl, O-alkenyl, chlorine, bromine, iodine, NO₂, amino, loweralkylamino, or di(loweralkyl)amino;

R⁸ is H, alkyl (including lower alkyl), chlorine, bromine or iodine; alternatively, R⁷ and R⁹, or R⁸ and R⁹ can come together to form a bond; and X is O, S, SO₂ or CH₂.

166. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

167. A use of a compound of the structure:

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168. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

169. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

170. A use of a compound of the structure:

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171. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

172. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

173. A use of a compound of the structure:

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174. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

175. A use of a compound of the structure:

or a pharmaceutically acceptable salt thereof, in the manufacture of a medicament for the treatment or prophylaxis of a host infected with the Hepatitis C virus.

176. A use of a compound of the structure:

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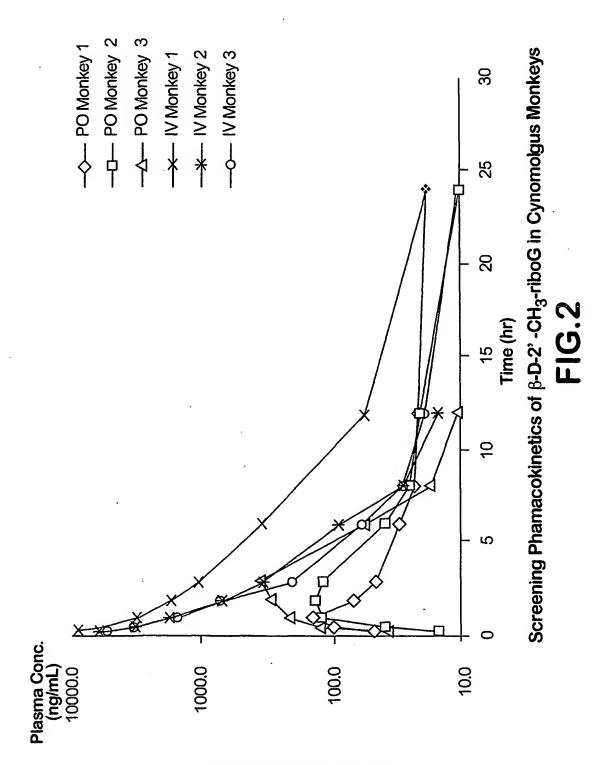
HO H₃C

- 178. Use of the compound as described in any of the preceding claims 130-177, wherein the said compound is in the form of a dosage unit.
- 179. Use of the compound of claim 101, wherein the dosage unit contains 178 to 1500 mg of said compound.
- 180. Use of the compound of claim 178 or 179, wherein said dosage unit is a tablet or capsule.

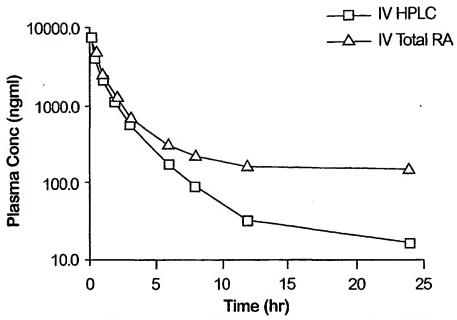
Chemical Structure of Illustrative Nucleosides

FIG.1

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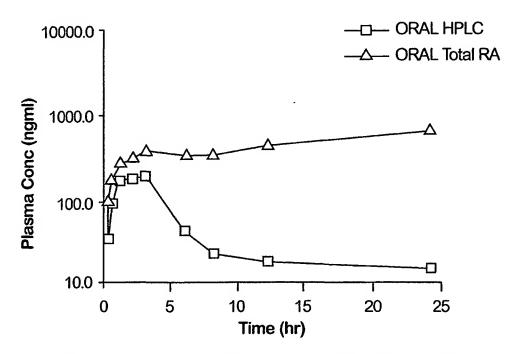


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Screening Phamacokinetics of β -D-2'-CH $_3$ -riboG in Cynomolgus Monkeys

FIG.3a



Screening Phamacokinetics of $\beta\text{-D-2'}$ -CH $_3\text{-riboG}$ in Cynomolgus Monkeys

FIG.3b

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 Peachtree Steet. Atlanta. GA 30303-1763 (US).

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In. ational Application No PCT/US 01/16671

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C07H19/06 C07H C07H19/10 C07H19/16 C07H19/20 A61K31/7068 A61K31/7076 A61P31/14 According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C07H A61K A61P IPC 7 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, PAJ, CHEM ABS Data C. DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. Υ WO 99 43691 A (CHOI YONGSEOK ; CHU CHUNG K 28-39. (US); HONG JOON H (US); SHI JUNXING (US) 52-63. 2 September 1999 (1999-09-02) 76, 79-90. 103-114, 127, 130-141, 154-165, 178 compounds 30,31 page 11, lines 25-31 the whole document -/--X Further documents are listed in the continuation of box C. Patent family members are listed in annex. Special categories of cited documents : "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the "A" document defining the general state of the art which is not considered to be of particular relevance invention "E" earlier document but published on or after the international "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to filing date "t" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "&" document member of the same patent family Date of mailing of the international search report Date of the actual completion of the international search 04. 03 2002 6 February 2002 Name and mailing address of the ISA Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl. de Nooy, A Fax: (+31-70) 340-3016

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| VIRUSES: MOLECULAR IDENTIFICATION AND TARGETED ANTIVIRAL THERAPIES" PROCEEDINGS OF THE ASSOCIATION OF AMERICAN PHYSICIANS, BLACKWELL SCIENCE, INC, CAMBRIDGE, MA, US, vol. 110, no. 2, 1998, pages 98-112, XP000885891 ISSN: 1081-650X | 130 |
| | , 103 |
| | |

international application No. PCT/US 01/16671

INTERNATIONAL SEARCH REPORT

| Box I | Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet) |
|-----------|--|
| This Inte | ernational Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons: |
| | Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely: |
| | Although claims 79-129 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound. |
| | Claims Nos.: bseaus URJ HERE to NEORMAT I ON not been applicated to the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically: |
| | |
| 3. | Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a). |
| Box II | Observations where unity of invention is lacking (Continuation of item 2 of first sheet) |
| This Inte | emational Searching Authority found multiple inventions in this international application, as follows: see additional sheet |
| | |
| | |
| 1 | As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims. |
| 2. | As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee. |
| 3. | As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.: |
| | |
| 4. | No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: |
| | |
| Remark | on Protest The additional search fees were accompanied by the applicant's protest. |
| | No protest accompanied the payment of additional search fees. |

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 7-12, 25-27, 34-39, 58-63, 76-78, 85-90, 109-114, 127-129, 136-141, 160-165, 178-180 (all partially)

The initial phase of the search revealed a very large number of documents relevant to the issue of novelty. So many documents were retrieved that it is impossible to determine which parts of the claims may be said to define subject-matter for which protection might legitimately be sought (Article 6 PCT). For these reasons it appears impossible to execute a meaningful search and/or to issue a complete search report over the whole breadth of the above mentioned claims. Consequently, the search has been restricted to the compounds of the above mentioned claims where R6 is methyl, ethyl, propyl, butyl, CF3 or Br-vinyl. Furthermore, in the case where R6 is methyl for compounds XI, XIV, XVII, or XVIII of the above mentioned claims, only several documents were cited.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. Claims: 1,4,13-18,25-27 (in part),28,31,40-45,52,55,64-69,
76-78 (in part),79,82,91-96,103,106,115-120,
127-129 (in part),130,133,142-147,154,157,166-171,
178 (in part),180 (in part)

Compounds of Formula I of claim 1 or compounds of Formula IV of claim 4, pharmaceutical compositions and uses pertaining thereto.

2. Claims: 2,5,19-24,25-27 (in part),29,32,46-51,53,56,70-75, 76-78 (in part),80,83,97-102,104,107,121-126, 127-129 (in part),131,134,148-153,155,158,172-177, 178 (in part),179, 180 (in part)

Compounds of Formula II of claim 2 or compounds of Formula V of claim 5, pharmaceutical compositions and uses pertaining thereto.

3. Claims: 3,6,25-27 (in part),30,33,54,57,76-78 (in part),81, 84,105,108,127-129 (in part),132,135,156,159, 178 (in part),180 (in part)

Compounds of Formula III of claim 3 or compounds of Formula VI of claim 6, pharmaceutical compositions and uses pertaining thereto.

4. Claims: 7,25-27 (in part),34,58,76-78 (in part),85,109, 127-129 (in part),136,160,178 (in part), 180 (in part)

Compounds of Formulae VII or VIII or IX of claim 7, pharmaceutical compositions and uses pertaining thereto, where the compounds do not fall within one of the earlier described subjects.

5. Claims: 8,25-27 (in part),35,59,76-78 (in part),86,110, 127-129 (in part),137,161,178 (in part), 180 (in part)

Compounds of Formulae X or XI or XII of claim 8, pharmaceutical compositions and uses pertaining thereto, where the compounds do not fall within one of the earlier described subjects.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

6. Claims: 9,25-27 (in part),36,60,76-78 (in part),87,111, 127-129 (in part),138,162,178 (in part), 180 (in part)

Compounds of Formulae XIII or XIV or XV of claim 9, pharmaceutical compositions and uses pertaining thereto, where the compounds do not fall within one of the earlier described subjects.

7. Claims: 10, 25-27 (in part),37,61,76-78 (in part),88,112, 127-129 (in part),139,163,178 (in part), 180 (in part)

Compounds of Formula XVI of claim 10, pharmaceutical compositions and uses pertaining thereto, where the compounds do not fall within one of the earlier described subjects.

8. Claims: 11,25-27 (in part),38,62,76-78 (in part),89,113, 127-129 (in part),140,164,178 (in part), 180 (in part)

Compounds of Formula XVII of claim 11, pharmaceutical compositions and uses pertaining thereto, where the compounds do not fall within one of the earlier described subjects.

9. Claims: 12,25-27 (in part),39,63,76-78 (in part),90,114, 127-129 (in part),141,165,178 (in part), 180 (in part)

Compounds of Formula XVIII of claim 12, pharmaceutical compositions and uses pertaining thereto, where the compounds do not fall within one of the earlier described subjects.

information on patent family members

PCT/US 01/16671

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